THE CHEMICAL FORMULARY

A CONDENSED COLLECTION OF VALUABLE, TIMELY,
PRACTICAL FORMULAE FOR MAKING THOUSANDS
OF PRODUCTS IN ALL FIELDS OF INDUSTRY

VOLUME III

Editor-in-Chief

H. BENNETT





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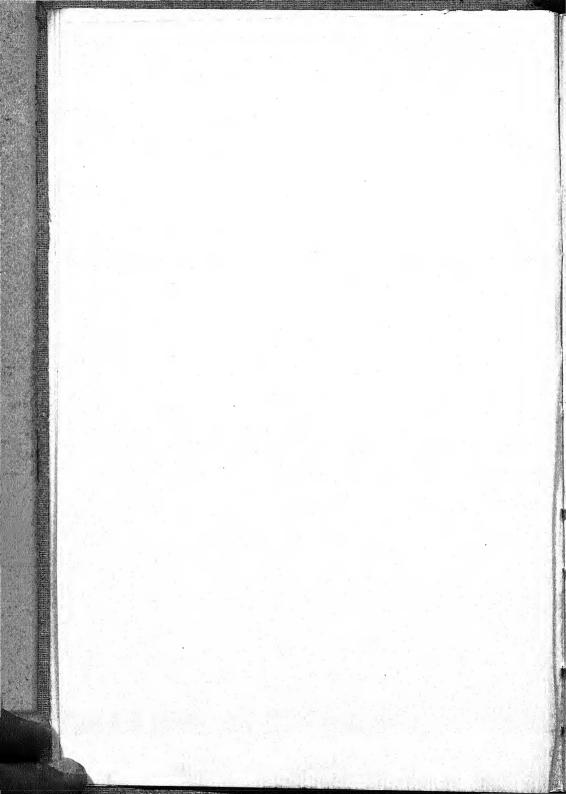
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PREFACE TO VOLUME II

The gratifying reception accorded Volume I of the Chemical Formulary together with the helpful and constructive criticisms received from reviewers and chemists have manifestly proved the need for a book of this type covering modern formulation in commercial chemistry.

While Volume I is complete in itself, the Editors felt it was impossible within the scope of one book to include all the formulae compiled for the numerous subject headings in the book. Volume II therefore is not a duplication or revision of Volume I but an entirely new work giving further formulae on the subjects treated in the first volume as well as more detailed information on processes and fundamental principles involved.

It will be noticed that all patented formulae have the patent number included. A helpful article on what is patentable in chemical compounding: infringements, licensing, etc., is another important addition to the book. It must be borne in mind in this connection that patented formulae cannot be used in the manufacture of commercial products unless prior arrangements have been made with the patentee.

The Editorial Board has been considerably enlarged and consequently it has been possible to include formulae hitherto unavailable.

A certain amount of criticism was directed toward the use of trade-names in Volume I. It was contended by the critics that formulae containing trade-names should be eliminated regardless of their value. Considerable thought was given to this contention and it was felt that, inasmuch as chemical trade-name products are being used in an ever-increasing number of formulae in every class of chemical manufacturing, these formulae should be included unless the application was exceptionally limited.

A second subject of criticism was the non-uniformity of systems of weights and measures used in the book. Since there is no uniformity in such systems in commercial practice and since the main purpose of the book is to familiarize the reader with commercial practice it was thought best not to attempt to standardize these systems.

In the Preface to Volume I, it was emphasized that the chemistry taught in schools and colleges is rightly confined to synthesis, analysis and engineering whereas in commercial manufacture many of the products so made are not synthetic or definite chemical materials but consist of mixtures, blends or highly complex compounds.

Because of the paucity or antiquity of the literature in this field and because of the difficulty encountered even by experienced chemists on entering new fields a definite need has existed for a modern compilation of formulae for chemical compounding and treatment.

In addition to an Editorial Board composed of chemists and engineers in many industries, publications, laboratories, manufacturers and individuals have been consulted to obtain the latest and best information in the numerous fields covered in the book.

It is important to remember that repeated experiments may be necessary to get the best results, especially when the field is intricate or unfamiliar. Again, although many of the formulae are being used commercially, some of them have been taken from patent specifications and the literature. Since these sources are subject to various errors and omissions, due regard must be given to this factor.

Formulae must be considered chiefly as starting points, variations have to be made to meet individual requirements and specifications. In cases of doubt or difficulty it is advisable at all times to consult other chemists or technical workers familiar with the particular field. This applies particularly in the case of the layman, as while a certain expense is involved this is more than compensated for by the saving of time, money and material.

As mentioned in Volume I it is hoped that those who have found a work of this kind helpful, will bring to our attention any errors they come across and will fee free at all times to make any constructive criticisms or suggestions.

PREFACE TO VOLUME III

Because of an insistent demand for new and additional formulae Volume III of the Chemical Formulary is being published a year in advance of original plans. In technical chemical compounding there is no rest or "breathing-spell"—no "status quo." Improvements are being made daily and new ideas and methods are continually being initiated and applied. New sources of data in many fields are being opened up in order to increase the breadth and scope of information.

As far as possible there has been included information especially requested by users of Volumes I and II. Diligent cooperation on the part of many chemists, engineers, teachers, technicians and other workers has made this possible.

The editor-in-chief wishes to thank all those who have helped in this work, which, in so short a time, has found a place as a highly useful tool and time-saver at the right hand of so many technical workers. In many cases it has proved to be a veritable catalyst in stimulating new products and processes.

Any thoughts for improving succeeding volumes and any new formulae or data, will, as heretofore, be most welcome. To make reference more easy the index in this volume is inclusive of Volumes I, II and III so that three separate indices need not be consulted.

H. BENNETT



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ABBREVIATIONS

amp
b.pboiling point
Be Baumé
CCentigrade
C Centigrade cc cubic centimeter
c.dcurrent density
c.pchemically pure
c.d. current density c.p. chemically pure cu. in. cubic inch
cu. itcubic foot
ddensity
dildilute
dr dram
F Fahrenheit
I.I.Cfree from chlorine
f.f.p.a free from prussic acid
fl. drfluid dram
fl. ozfluid ounce
ggram
grgrain
hrhour
kgkilogram
lliter
m.pmelting point
minminute
minminims
NNormal
pH
Q. S A quantity sufficient to make
r.p.mrevolutions per minute
sec
sp. Gspecific gravity
rich din square decimeter
U.S.P U. S. Pharmacopeia
Vvoltage
wt. ,,,,,,,,,,,,weight

ADHESIVES

1/4 lb.

White Glue

A solution consisting of:	
Animal Glue	100 oz.
Zinc Oxide	50 oz.
Water	100 oz.
gives a glue which sets quite very strong.	hard and is

Glue
Urea 1 lb.
Casein 2 lb.

Hydrate of Lime

Black Albumen from Blood

Let slaughterhouse-blood stand in shallow dishes or pans, cut the blood jelly, sift the serum off. The residue is stirred in water to a paste, and put through a filter press. Evaporation in vacuum produces from the second filtrate the dark black albumen used for veneering and laminating.

"Salamyn-Plant" Glue

	Switching in a recur	o arac	
a.	Potato-Starch	35	
	Water (35° C.)	105	l. •
	Caustic Soda (35°		kg.
c.	Hydrochloric Acid	about 10	kg.
	Water	10	l.
d.	Upholsterer's Glue	260	kg.

Stir a for $\frac{3}{4}$ hour after adding d. Stir with b until glassy, then add c.

Calcium Saccharate Glue

Water, Boiling	70 g.
Sugar	6 g.
Lime, Fresh Slaked	1.5 g.

Let stand, stir often, cover. After a few days pour off from bottom deposit, and soak in the solution,

Carpenter's Glue 60 g. then warm to solution.

Marine Linoleum Cement

Decks to be covered with linoleum should be thoroughly cleaned, and the linoleum stuck to the deck with the following adhesive:

To make 10 gallons, first cut 4 oz. of crude (ham) rubber into small lumps and

dissolve in 4½ gallons of gasoline. It will require about two days to get the rubber into colloidal solution. When in proper condition it should string about two inches thumb and forefinger. Cut 19 lb. of gum shellac in 34 lb. of denatured (or wood) alcohol. Add 62 lb. of whiting then add the rubber solution. For best results this mixture should be ground in an iron or pebble mill.

Linoleum Glue

a. Rye or Barley Flour	50 kg.
Water, tepid	250 1.
b. Caustic Soda (20° Bé.)	20 kg.
c Turnentine Venice	-

melted 20-25 kg. Part a dispersed by stirrer is mixed with b (dissolved). The mixture is then boiled, and after cooling emulsified by adding c (while stirring add).

Painters' Glue (Cold)

Water (25° C.)	350 1.
Potato-Starch, Powder	100 kg.
Rosin, Finely Ground	21 kg.
Caustic Soda (24° Bé.)	56 kg.

Mix altogether with strong stirring for 2-3 hours, let stand 1 hour, and neutralize with dilute nitric acid until red color with phenolphthalein disappears (in a sample). Stir ½ hour more.

Wall Size

Aluminum Stearate		4	oz.
Turpentine		25	
Mineral Spirits (150-190°	C.)	71	oz.

Heat the turpentine to 180° F. and add the stearate slowly while stirring continuously. Add mineral spirits and stir until clear.

Painters' Size

Potato-Starch (Air-Dried)	7.8 g.
Calcium Chloride	7.0 g.
Water	3.0 g.

The aqueous paste, when compact, is dried and ground. The excess chloride can be extracted with aqueous alcohol, yielding a better paintable and quicker drying product.

Paperhanger's Paste

Use a cheap grade of rye or wheat flour, mix thoroughly with cold water to about the consistency of dough or a little thinner, being careful to remove all lumps. Stir in a tablespoonful of powdered alum to a quart of flour, then pour in boiling water, stirring rapidly until the flour is thoroughly cooked. Let this cool before using and thin with cold water.

Venetian Paste

a.	White or Fish Glue	4		oz.	
	Cold Water	8		OZ.	
	Venice Turpentine	2	fl.	OZ.	
c.	Rye Flour	1		1b.	
	Cold Water	16	fl.	oz.	
d.	Boiling Water	64	fl.	OZ.	

Soak the 4 oz. of glue in the cold water for 4 hours. Dissolve on a waterbath (glue-pot) and while hot stir in the Venice turpentine. Make up c into a batter free from lumps and pour into d. Stir briskly, and finally add the glue solution. This makes a very strong paste, and it will adhere to a painted surface, owing to the Venice turpentine in its composition.

Flour Paste

a.	Wheat Flour	2	lb.	
	Cold Water (1 quart)	32 f	l. oz.	
b.	Alum	1	oz.	
	Hot Water	4 f	l. oz.	
c.	Boiling Water	96 f	l. oz.	

Work the wheat flour into a batter free from lumps with the cold water. Dissolve the alum as designated in b. Now stir in a and c and, if necessary, continue boiling until the paste thickens into a semi-transparent mucilage, after which stir in the solution b. This makes a very fine paste for wallpaper.

Sinclair's Glue

Formula No. 1

"Very Good" Glue or			
Gelatin		50	oz.
Water		100	oz.
Glycerin	4	or 6	oz.
Thymol or Menthol		0.15	oz.

The smaller amount of glycerin is for summer or tropical use, and the larger amount for winter. Gelatin is preferable, for commercial glue varies in quality and generally requires neutralizing to litmus with weak alkali. The following is a simple test for a "very good" glue. "On soaking glue in excess of cold

water overnight, a gelatinous coherent mass is obtained, weighing, when drained, at least four times the weight of the original glue.'' With the very best glue a mass weighing five times the original weight is obtained.

No. 2	2			
Isinglass			5 0	oz.
Gelatin			50	OZ.
Water			200	CZ.
Tannic Acid			12	OZ.
Glycerin	8	or	more	oz.
Menthol or Thymol			0.15	OZ.

This forms a stronger adhesive, is perhaps more elastic, and has the advantage of somewhat hardening the skin so that the tendency to blistering is almost completely eliminated.

Marine Glue .

Rubber	100	g.
Turpentine	600	g.
Coal Tar Oil	600	g.
Shellac	300	ğ.

Warm together and mix till smooth.

Preserving Glue

Add 3 ounces of ordinary borax to each gallon of glue or add 1 ounce of formaldehyde to the gallon or 1 ounce of carbolic acid. Adding ½ ounce of 28% acetic acid to 2 pounds of glue will also prevent the souring and also has a tendency to make it waterproof.

Casein Glue

Formula No. 1

Casein			100	oz.
Water	220	to	230	oz.
Hydrated Lime	20	to	30	oz.
Water			100	oz.
Silicate of Soda			70	oz.
Copper Chloride	2	to	3	oz.
Water	30	to	- 50	oz.

The 220 to 230 parts of water added to the casein is approximately the right amount to use with Argentine (naturally soured) casein; but if a different casein is used the water requirement will lie somewhere between 150 and 250 parts by weight. The correct amount for different caseins must be determined by trial.

The formula presupposes that a high calcium lime will be used. A lime of lower grade may be used, but a proportionately larger amount of it will be needed, or the water resistance of the glue will be sacrificed. It is suggested that for the first trial the user try 25 parts of lime. If this does not give

proper results the amount can be varied within the limits specified.

The density of the silicate of soda used should be about 40° Baumé, with a silica-soda ratio of from 3 to 3.25.

Copper sulphate can be substituted for

copper chloride.

Place the casein and water in the bowl of a mixing machine and rotate the paddle slowly, stirring the mixture until all the water has been absorbed and all the casein moistened. If the casein is allowed to soak beforehand it is more readily dissolved in the mixing process. Mix the hydrated lime with water in a separate container. Stir this mixture vigorously at first, but just before it is added to the casein stir just enough with a gentle rotary motion to keep the lime in suspension. Pour the milk of lime quickly into the casein.

When casein and lime are first combined they form large, slimy lumps, which are balls of dry casein coated with partly dissolved casein. These break uprapidly, becoming smaller and smaller, and finally disappear. The solution, in the meantime, is becoming thin and fluid. At this point stop the paddle and scrape the sides and bottom of the container, and then stir again. If a deposit of casein remains unacted on, it may cause

more lumps later.

When about two minutes have elapsed since the lime and casein were united, it may be noticed that the glue has begun to thicken a little. Add the sodium silicate now, or else the glue will become too thick. The glue will momentarily become even thicker, but this thickness will soon change to a smooth and fluid consistency.

Continue the stirring until the glue is free from lumps. This should not take more than 15 or 20 minutes from the time the lime was added. If the glue is a little too thick, add a small amount of water. If the glue is too thin, it will be necessary to start over again, using a

smaller proportion of water.

The copper salt may be added at any one of several times during the mixing process. If added as a powder before the casein is soaked, it may have a corrosive action upon the metal container. The copper salt, if added as a powder, should be thoroughly mixed with the casein before the addition of the lime. Copper salt may be placed in solution and conveniently stirred into the moistened casein immediately before the lime is added or after all the other ingredients have been combined. If the copper solution is added at the end of the

mixing period, pour it into the glue in a thin stream and stir the mixture vigorously. Continue stirring until any lumps, which may have formed by the coagulation of the glue and the copper solution, are broken up and until a smooth violet-colored glue is obtained.

smooth violet-colored glue is obtained.

Glue prepared by this formula has proved to be exceptionally strong and durable, even under wet or damp con-

Formula No. 2

The mixing is the same as for above formula except for the omission of the copper chloride. The glue made by this formula has a medium consistency, excellent working properties, a good working life, and makes joints of high strength, but it falls somewhat short of the previous formula in water-resisting properties, especially when the lower amounts of lime are used.

Casein Water	100 200	
Sodium Hydroxide (Ca Soda) Water	10	oz.

Bring the casein and water together according to the directions for mixing glue prepared by previous formula. Dissolve the caustic soda in water in a separate container, and while the mixing paddle is revolving sprinkle the caustic soda solution into the damp casein. Str slowly until a thin, smooth glue has been obtained. The consistency of the finished product may be altered by adding more casein if it is too thin, or by adding water if it is too thick. Silicate of soda is sometimes added to thicken or to reduce the cost of the glue per unit of volume.

This glue has exceptional strength when dry, but when exposed to moisture it weakens as rapidly as animal or vegetable glue.

Cold Glue (Casein)

Formula No. a. Casein, Dry Trisodium Phosphate	1	70 10	ğ.
Lime Hydrate Sodium Fluoride b. Water		200	g. g.
Pine Oil a is soaked with b.		2	g.

	No. 2	
a.	Casein	60 g.
	Lime, Hydrated	15 g.
	Trisodium Phosphate	4 g.

F.			THE C	HEMICA
n	Sodium Fluc Nut Meal b. Water Stir a with by		ready	4 g. 17 g. 200 g. after 20
	Casein Caustic Soda (36° Bé.)	No. 3		-30 g. -0.6 g. 7-2 g.
	Water Casein	No. 4		–68 ec. –30 g.
	Soda Ash Water	No. 5	2 78-6	4.5 g. 35.5 cc.
	Casein Borax Water	No. 6	78	–30 g. – 5 g. –65 cc.
	Casein Ammonia (sp. Water	gr. 0.93	10) 10	–30 g. –24 ec. –46 ec.
	Casein Borax Ammonia (0.91 Water		12 g. 1.5 g. 1.5 g. 85 g.	Knead
	Casein Water		soak	
	Disodium Hy- drogen Phos- phate Water Caustic Soda (10%)		dissolv	Knead
	Mix all in war	m water No. 9	-bath.	
	Casein Water Borax Ammonia (0.9 Caustic Soda (36° Bé.)	1)	20 g. 80 g. 1 g. 2 g. 2 g.	Warm for ½–1 hour
	Cool, at 50-60 Waterglass (3 Alcohol, Dena	° C., ad 0° Bé.) tured	ld:	8 g. 2 g.
	Impre Casein Ammonia (sp. Water	egnation g. 0.91	0) 1	5–20 g. 3–16 cc. 7–64 cc.
	Casein Ammonia (0.9	astel''(10)	Glue	25 g. 20 cc.

Water		50	cc.
Glue Jelly	. 1	5	g.

Modern Casein Adhesive Powders

For use stir with 140 times the amount of water (cold). After ½ to ¾ hour, a homogeneous, viscous solution is gotten ready for use.

Formula No. 1		
Lactic Acid-Casein Marble-Lime Hydrate Trisodium Phosphate Sodium Fluoride Sodium Sulphate, Pure,	70 13 5 4	ත් ක් ත් ක්
Anhydrous Naphtha, Refined	$\frac{6}{2}$	g. g.
Lactic Acid-Casein Slaked Lime Trisodium Phosphate Aniline Mineral Oil No. 3	60 20 10 8 2	ත් ත් ත් ත් ත්
Lactic Acid-Casein Slaked Lime Trisodium Phosphate Sodium Sulphite Sodium-Waterglass, Dry Copper Chloride Hardwood-Meal Mineral Oil	50 16 8 8 6 2 10 11/2	දුරු දුරු දුරු දුරු දුරු දුරු දුරු

Air-plane	Prop	elle	r Glue
1. Black Blood Albumen Water	1 6	g. }	Mix at 15° C., stop mixing for two hours
Add: Slaked Lime Water	0.06 1	$\left. egin{matrix} g. \\ g. \end{matrix} \right\}$	Mix until thick

Mordant for Handles of Kitchen-Knives

a. Potassium Bicaromate 15 g.
Water 1000 cc.
b. Ammonia (25%) 150-200 g.

Dissolve the chromate a, and add b.

Treat wood with solution, dry, rub over with a hard brush (horse-hair), optionally a thin polish.

Wood Veneer Adhesive U. S. Patent 1,964,960

Casein 1 oz.

Ammonium Sulphocyanate 2 oz.

Paraformaldehyde .02–0.4 oz.

Water sufficient to make fluid.

This will remain fluid for several hours at ordinary temperature. Coagulates on heating to give strong bond.

Cement for Filling Cracks in Wood Consists of a mixture of 150 parts linseed oil, 30 parts varnish, 40 parts wax, 30 parts gypsum, 750 parts pigment.

(Note: Generally, wax is an objectionable constituent, from the standpoints of lessening adherence within the crevices and lack of cohesion of finishing coatings applied over such filled areas. Preferable material would be the present well-known plastic wood and wood doughs which are pyroxylin-base products utilizing wood flour. Representative composition (U. S. Patent 1,838,618) is Celluloid Scrap 19 lbs., Ester Gum 8 lbs., Castor Oil 3 lbs., Methyl Acetone 44 lbs., Wood Flour 26 lbs.; and if pigmentation may be desired, as follows—Celluloid 10, Ester 7, Castor 4, Acetone 15, Benzol 15, Alcohol 5, Wood Flour 24, China Clay 20.

Cheaper materials more popular with painters and decorators are the Water Putties in dry powder form; they are used for filling cracks and holes in wood trim, also for filling the spaces between flooring in both old and new floors. When thoroughly dry the applied putty has no tendency to shrink or crack. One product on the market for years is composed of 10 parts Quartz Silica, 2 parts Plaster of Paris, 11/2 parts Dextrin. Pulverized Gum Arabic could be substituted for the dextrine and effect greater hardness; and addition of about one-half part of wood flour or fine sawdust would enhance the toughness of the putty. For using, only enough water is mixed with the putty powder to the consistency of regular commercial putty).

Wood Veneer Glue

Blood Albumen		40	g.
Casein		12	
Slaked Lime	-	6	g.
Sodium Fluosilicate		2	g.
Wood Meal		40	g.

Apply the adhesive by putting it on both sides of the middle piece of wood. If the adhesive is just too viscous, homogenize the adhesive layer. The wood pieces are put together, then pass through drying chambers at 90-95° C., under a pressure of 12-18 kg. per cc. until the albumen is coagulated.

Sealing Preparation for Wine-Barrels Vaseline (40–42° C.) or socalled 'Traction-Paraffine' (42-44° C.) 98-98.5 g. Tallow, Hard Fat or Palm O53 2- 1.5 g.

Impregnating "Green" Wood Austrian Patent 142,431

Cover with the following paste and allow to remain until dry.

~ 1 77 17	80 lb.
Sodium Fluoride	
Sodium Dinitrophenolate	15 lb.
Kieselguhr	5 lb.
Water sufficient to make pas	ste.

Gum Arabic Glue

Gum Arabic Lime Water, Saturated	15-20 g. 10-20 cc.
Glycerin	1-3 g.
Water	74-27 cc.

Mucilage		
Gum Arabic, Amber Sorts	100	lb.
Water	150	lb.
Heat and stir until dissolved.		
Strain and add		
Oil of Cloves	5	oz.
Oil of Wintergreen	5	oz.
Salicylic Acid	5	oz.
Photo-Paste		
Gum Arabic	30	g.
Saturated Lime Water		cc.

Cold Water Paste Australian Patent 8259

10 cc.

45 cc.

2% Tragacanth Solution

Water

Wheat Flour		8	oz.
Alum		1	oz.
Water		8	oz.
Mix till smooth; powder.	evaporate	till	dry

Pasting Paper on Metal Surface

- 1. Clean off grease with hot soda solution.
- aper.

Ζ.	Kougnen	with	emery	р
	Prepare			_

٠	Frepare giue:	
	a. Water	4 kg.
	Calcium Chloride	1 kg.
	b. Bone Glue	1-2 kg.

Dissolve a, then swell b in the solution for 24-30 hours; heat on water bath to obtain solution.

Moldex or other preservative 0.1-0.2%.

Vegetable Mucilage

- a. Water (Above 16° C.) 200 1. Potato-Starch 100 kg. b. Caustic Soda (35° Bé.) 28 kg.
- Stir a to dispersion, sift, add slowly b under stirring, until glassy. Keep temperature low if thick mucilage is wished.

Higher temperature yields more fluid glues.

Library Adhesive Paste

a. Capillary Syrup		
(42–44° Bé.)	70	kg.
Water, Boiling	20	1.
b. Water, Boiling Borax	8	kg.
	2-3	kg.
d. Sulphurous Acid (5° Bé.)	0.5	kg.
e. Formalin	0.5	kg.

Add b, c, d, e, in the given order separately to a, stirring strongly. When ready, dye with a little burnt sugar color.

Carton Glue

Dextrin, Light Borax Solution	100	g.) dissolve
(10%)	70	g.	dissolve hot
Caustic Soda (40° Bé.)	5	g.	add when cool
Let stand several		Ū	,

Cardboard Glues

1. Casein	13 g.
Trisodium Phosphate	1 g.
Ammonia (0.91)	2 g.
Water	85 g.
2. Casein	10 g.
Borax	2 g.
Glucose	2 g.
Waterglass (30° Bé.) 15 g.
Water	71 g.

Padding Glue

1.	Glue (Nat. Assoc.				
	8-10 Grade)	10	lb.		
2.	Glycerin	10	lb.		
	. Water	12	lb.	2	oz.
4.	Zinc Oxide	1	lb.	3	oz.
5.	. Beta Naphthol			1/4	oz.
6.	. Methyl Salicylate			1	oz.
. 7/	Tiv 2 and 4 thon	44	5 0:	nd s	2 an

Mix 2 and 4, then add 5 and 3, and then 1. Let stand over night, warm and stir until uniform; add 6 and pack.

In hot humid weather this glue may set too slowly. This may be corrected by

a. Using a higher grade of glue, or
 b. Using less glycerin (which will, of course, reduce flexibility), or

c. Dusting surface after partial drying with talc or precipitated chalk.

Tabbing Compound U. S. Patent 1,966,389

775 parts of uncoagulated vulcanized latex, containing 40 to 42% by weight of

total solids constitutes the first ingredient.

The second ingredient is prepared by dissolving 50 parts of casein in about 150 parts of distilled water (preferably with the addition of an alkali which may be caustic soda, alkaline sodium salts or ammonia).

Third, 50 parts of egg albumen are dissolved in about 200 parts of water to produce a highly viscous solution.

A fourth component is made by adding 125 parts of a 2% ammonia solution, to 5 parts of dried wood fibre and 5 parts of cellulose flocks (or other fibrous material) and the mixture is stirred until a substantially uniform suspension is produced. A small amount of a deodorant composition such as oil of wintergreen can also be added at this point if desired.

The casein solution and the egg albumen solution are then added slowly with constant stirring to the vulcanized uncoagulated latex, and the stirring is continued until a uniform or homogeneous mass is produced. If desired, suitable coloring materials can be added at this stage and can be thoroughly stirred in.

The ammoniacal liquor containing the fibrous material "fourth component" is then added and the entire mixture thoroughly stirred or churned in order to produce a uniform mixture. This mixture is then ready to be used for tabbing, or it can be simply canned and used at any subsequent time.

For tabbing, the paper is jogged if desired to give a substantially smooth surface of edges, to which one coat of the material is brushed on rapidly. after five or ten minutes a second coat is preferably applied. This second coat can be daubed on heavily, and quickly brushed down to a smooth coating. The composition will dry firm and the exposed surface will be substantially free from tackiness in about half an hour or sometimes twenty or twenty-five minutes, depending upon atmospheric conditions. The complete strength of the composition is however not developed for several hours after application. If desired, the tablets can be allowed to stand quiet for several hours, until substantially the maximum strength has developed. The surface can be finally dusted over with a suitable pulverulent material, such as talc powder if desired, although ordinarily this will not be found necessary, since the composition after drying does not stick to other surfaces with which it comes into contact, at least to an objectionable degree.

oz.

The brushes or the like used in applying the composition can be readily cleaned by being washed in water, and any of the material which gets onto the hands of the user can be readily washed off with water.

In case the solution becomes too thick, it can be diluted with soft water (preferably rain water or distilled water). Hard water would be injurious to the compound.

Label Gum

Label Gum		
Formula No. 1—Fluid		
Gum Arabic Saturated Lime Water Glycerin Water No. 2—Less Fluid	$30 \\ 15 \\ 1 \\ 54$	g. ec. g. ec.
Gum Arabic Aluminum Sulphate Crystals Glycerin Water No. 3—Viscous	$\begin{array}{c} 35 \\ 2 \\ 2 \\ 61 \end{array}$	g. g. g. c.
Gum Arabic Aluminum Sulphate Crystals 2% Tragacanth Solution Water	$\begin{array}{c} 30 \\ 2 \\ 20 \\ 48 \end{array}$	
Label Glue		
Formula No. 1 Casein Ammonia (sp. g. 0.910) 30% Rosin Soap Water	20 16 5 59	g. ec. g. ec.
No. 2		
Water-Resistant Casein Ammonia (0.910) Waterglass (30° Bé.) Water	20 5 6 70	oz.
Library Mucilage		
Formula No. 1—Fluid		
Gum Arabic Saturated Lime Water Glycerin Water	$25 \\ 15 \\ 1 \\ 59$	g. cc. g. cc.
No. 2—Less Fluid		
Gum Arabic Lime Water, Saturated Glycerin Water	$\frac{40}{20}$ $\frac{2}{38}$	g. cc. g. cc.
No. 3—Viscous	00	
Gum Arabic Aluminum Sulphate Crystals 2% Tragacanth Solution	20 2 15	g. g. cc.

63 cc.

Water

	F	aper M	ucilage	
a.	Dextrin,	Middle	Pale	5

Water 50 oz.
b. Sodium Bisulphite 0.5 oz.
Borax 1.0 oz.
Camphor a grain

Stir cold until lump-free, warm until the mucilage is formed. Add b for deodorizing and preservation.

Adhesive for "Gumming"	Papers
Gum Arabic	30 g.
Saturated Lime Water	15 cc.
Glycerin	2 g.
2% Tragacanth Solution	5 cc.
Water	48 cc.

	Paper Bag Glue	
Casein		22 g.
Borax		3 g.
Venice	Turpentine	3 g.
Water	_	72 cc.

The casein has to be treated (swelled) at 50-70° C. When treating with ammonia, heat up higher at the end to evaporate the excess. Moldex or other good preservative is to be added after the alkaline treatment in proportions of about 18-25 ounces per 100 gallons. If too viscous or too thin, add or evaporate water.

Let stand to clear up.

Carton Glue

Casein	25	g.
Caustic Soda (36° Bé.)	0.5 or 1.7	
30% Rosin Soap	10	g.
Water	64.5 - 63.3	cc.

Waterproof Adhesive U. S. Patent 1,965,778

Formula No. 1 Casein 100 lb. Water 225 lb. *Wax Solution 3 lb. No. 2 Vegetable Protein Glue 100 lb. Water 325 lb.

Vegetable Protein Glue	100 lb.
Water	325 lb.
*Wax Solution	3 lb.
* Consists of: Carbon Bisulphide	0.15
Carbon Tetrachloride	8 lb. 8 lb.
Paraffin Wax	1 lb.

Non-Caking Dextrin Adhesive French Patent 783,963

Dry adhesives having a basis of dextrin which dissolve in cold water without caking are made by heating dextrin to about 80° C. for 1/2 hour with about 1% of a polyhydric alcohol, e.g., glycol.

Mucilage for Paper, Photos, Printed Matter

a.	Soft Water	35 g.
	Sugar	1 g.
	Wheat Starch	3 g.
7.77	ann and stir until glassy.	

b. 19 parts of a 20-25% gum arabic solution.

Solution b is added to a when the starch has become "glassy." Preserve with phenol or oil of cloves.

Gummed Labels for Brass, Tin

O CTTTTTTT O TO		
Moisten with: Acetic Acid Glycerin Water	8 fl. 2 fl. 6 fl.	٥z.

U. S. Postage Stamp Glue

Gum Arabic	1	lb.
Starch	_	lb.
Sugar	4	lb.

Distilled Water sufficient to give desired consistency.

Adhesive for Waxed Papers

Formula No. 1 Thickened Spirit-Lacquer

Acetyl Cellulose-Solution

	No. Z	
Rosin		60 g.
		10 g.
Mastic		20 g.
Sandarac		
Ether		5 g.
		75–100 g.
Alcohol	0	10 200 8.
	NT ~ 2	

No. 3 Chromium Gelatin

or Canada Balsam

No. 4

a. Cologne Glue (or	
Galatin)	100 g.
b. Acetic Acid, Dilute	200 g.
c Potassium Bichromate	5 g.

Soak a in b, then dissolve on steam bath, add c. No 5

	740. 0			
Alcohol			100	g.
Ether			5	g.
Rosin		(30-70	g.
			20	g.
Sandarac			10	
Mastic			10	₽.

Celluloid Cements

Formula No. 1	
Pyroxylin	200 g.
Camphor	40 g.
Gum Elemi	8 g.
Amyl Acetate	2600 cc.
Acetone	500 cc.
Methanol	400 cc.
No. 2	
C.HLeid Chemines	240 g.
Celluloid Shavings	8 g.
Gum Elemi	500 cc.
Acetone	1500 cc.
Methanol	1500 cc.
Amyl Acetate No. 3	1000 001
210. 0	160 g.
Pyroxylin	40 g.
Camphor	2100 cc.
Methanol	1400 cc.
Fusel Oil	280 cc.
Castor Oil	200 00
No. 4	40
Celluloid Shavings	40 g.
Gum Elemi	8 g.
Benzol	1000 cc.
Amyl Acetate	1000 cc.
Methanol	800 cc.
Acetone	600 cc.
No. 5	
Pyroxylin	150 g.
Camphor	40 g.
Methanol	2525 сс.
Amyl Acetate	1260 cc.

Cement for Safety "Movie" Films

The formula below was developed especially for safety films and acetate type of transparent sheeting.

"	oranghar orre series		
	Cellulose Acetate	4	oz.
	Tri-Phenyl Phosphate	2	oz.
	Acetone	60	oz.
	Di-Acetone Alcohol	9	oz.
		15	0%
	Benzol	10	-
	Methanol	10	UZ.

The cellulose acetate of high viscosity film quality is preferred. However, washed safety movie film free from the gelatin coating, or other source of reclaimed cellulose acetate may be used. Instead of tri-phenyl phosphate plasticizers of the toluene sulphonamid type such as the Santicizers may be used.

Movie Film Cement

This composition is effective on either the inflammable or safety type films. In using this cement it is preferable to scrape off the gelatin coating with a knife or steel wool.

Cellulose Nitrate	4	oz.
Tri-Cresyl Phosphate	2	OZ.
Ethyl Acetate	55	oz.

Butyl Acetate	14 oz.
Benzol	15 oz.
Methanol	10 oz.

The cellulose nitrate may consist of a good grade of high viscosity nitro-cotton or clean new celluloid scrap or nitrate movie film with the gelatin coatings removed. If new cellulose nitrate is not used, the tri-cresyl phosphate can be reduced about one-half. The solvents are mixed together in the above proportions by weight and the cellulose nitrate added.

Pyroxylin Cement

Celluloid Scrap	40	g.
Amyl Acetate	350	cc.
Wood Alcohol	100	cc.
Ethyl Alcohol, Denatured	50	cc.
Gum Elemi	15	g.

Methyl Cellulose Adhesive

Methyl Cellulose	1 lb.
Water	40-60 lb.
Warm together and sti	r until uniform.

"Cellophane" Adhesive U. S. Patent 1.972.448

Chlorinated Polyphenyl Resin (125° C. softening	
point)	62.5 lb.
Dibutyl Phthalate	5.4 lb.
Silica, Finely Ground	32.1 lb.

Cigarette Paper Adhesive

Formula No. 1		
Pectin Bone Glue, Liquid	54 13.5	0Z.
Bone Glue, Solid	13.5	0Ζ.
Dextrin	19	OZ.
No. 2		

60.5 oz.

oz.

Pectin

Rye Flour

Bone Glue,	Fluid 16.5	oz.
Bone Glue,	Solid 6.6	oz.
Dextrin	12.5	OZ.
Rye Flour	4.0	oz.
	No. 3	
Pectin	50	oz.
Bone Glue,	Solid 10	oz.
Dextrin	10	07

In the above formulae add sufficient water to make a mucilage of desired consistency.

Primer for Wall Paper Paste U. S. Patent 2,005,900

Sodium Silicate		50	oz.
Water			oz.
Copper Sulphate	(121/2% So-		
lution)		6	oz.

Mailing Tube Adhesive

Glue, Ground Animal	40 oz.
Water	54.7 oz.
Nitric Acid	5.0 oz.
Phenol	0.3 oz.

Sealing of "Transparit," "Helioglas," or "Cellophane" Packages

a. Methyl Acetate 80 cc. Ethyl Lactate 20 cc.

 Collodion-Wool or washed film-scrap, as much as necessary to give a viscous solution (like 30-31° glycerin)

"Cellophane Adhesive"

Arabic, Gum	16.5 oz.
Glycerin	20.5 oz.
Glyceryl Bori-borate	9.0 oz.
Formaldehyde	4.5 oz.

Cardboard and Nitrocellulose Sheet Cement

U. S. Patent, 1,969,477

Nitrocellulose	4.5 oz.
Camphor	1.0 oz.
Acetone	30.0 oz.
Ethyl Lactate	10.0 oz.
Xylol	55.0 oz.
Water	5.0 oz.

Liquid Sealing Wax French Patent 751,683

Turpentine	100 cc.
Shellac	150 g.
Zinc Oxide	30 g.
Methanol	25 cc.

Mix until free from lumps. This dries in air after applying.

Elastic Sealing Wax

Rubber	Latex	(60%)	_	165	oz.
Shellac				12	OZ.

Warm together with stirring until all moisture is driven off.

De Khotinsky Type Laboratory Cement Improved Type

Shellac, Flake	100	g.
*Plasticizing Solvent	15 to 30	

Heat the solvent to 120° C., and slowly stir in the shellac flakes. When the shellac is thoroughly dissolved and the mixture homogeneous, cool slightly until the mixture pours with difficulty. Immediately pour into long tin molds of about one-half inch square cross section which have previously been treated lightly with petrolatum.

*As a "plasticizing solvent" pine tar has been widely recommended, but is inferior, since the excessive amount of 60 to 100 grams is required. The oil distilled from white-pine tar over the range of 200° to 325° C. is much better, yielding a tougher cement. Wood creosote or similar mixtures of substances like guaiacol, cresol and other low-melting, high-boiling phenois may be used; also trimethylene glycol or other slightly oxygenated organic solvents of high boiling point. The range of 15 to 30 grams approximately covers the variations of hardness commonly desired.

"Boltwood Wax"

(For cementing physical instruments)

Shellac	40 g.
Rosin	72 g.
Venice Turpentine	8 g.
Beeswax	60 g.
Talc, Dry	16 g.
Tin Oxide, Dry	16 g.
n = 1, 1, 2,	

Melt the rosin, add the shellac. Heat to 200° C., add the Venice turpentine and beeswax. Heat the mixture strongly with stirring until it ignites spontaneously. Let it burn until the total mass has shrunk to about 40% of its original weight, then add the talc and tin oxide. This gives a tough, smooth, waxy cement more easily handled on certain delicate instruments than the de Khotinsky type cement.

Leather Sole Cement

Nitrocellulose		22.5	g.
Alcohol		22.5	g.
Benzol		31.1	g.
Ethyl Acetate		9.5	g.
Camphor		1.1	g.
Acetone Oil		0.09	g.
Castor Oil	•	0.09	
			_

Cement for Leather or Leather on Rubber

Gutta-Percha	21.6	oz.
Carbon Bisulphide	17.7	oz.
Benzene	2.9	oz.
Turpentine Oil	23.5	oz.
Asphalt	34.3	oz.

Leather Cement

Celluloid	11.9	oz.
Naphthalene	1.2	oz.
Acetone	67.1	oz.

Cement for Stone and Leather, Porcelain and Leather, Glass and Leather

Crude Rubber	9.1	oz.
Heavy Benzine	45.5	oz.
Japan Wax	13.6	oz.
Colophony	31.8	oz.

Concentrated Rubber Cement German Patent 599,405

a.	Caoutchouc		10 g.
	Benzol		90 g.
b.	Nitric Acid	(52.77%)	1 g.

a gives after 24 hours stirring a homogeneous paste, which is depolymerized by adding b. When paste is dissolved, stop reaction by adding barium carbonate. Treat then with antimony trichloride or phthalic acid.

Rubber Cement

(Will firmly fasten rubber to almost any substance)

India Rubber (finely			
chopped)	100	oz.	
Rosin	15	oz.	
Shellac	10	oz.	
Carbon Disulphide, sufficient	to di	ssolve	

Softening Hardened Shoe Adhesive German Patent 605,725

Cellulose nitrate adhesives used in shoe cements are softened by the following:

Pyroxylin	(1100	second)	62	oz.
Alcohol	•	,	26	oz.
Acetone			450	oz.
α-Propylen	e Oxid	e	225	OZ.

Shoe Repair Cement U. S. Patent 2,004,059

Crepe Rubber		6	lb.
Rosin		21/2	lb.
Accelerator		11/2	lb.
Benzene		15	gal

Porous Leather Sealer

Shellac	14	lb.
Rosin	1	lb.
Alcohol	5	gal.
Butyl Alcohol	1/4	gal,
Castor Oil	4 🙃	oz.

Leather Belt Cement

	a.	Glue, Hide	50	g.
7	b.	Water	200	

Soak a in b, pour excess water off, and melt the soaked a with:

,	Glycerin		2%
•		Bichromate	2%
	Potassium	Dichromate	2%

When cooled, pour into oiled metallic forms; pack the gelatinous product at once into grease-proof paper.

Apply on roughed surface, while the sharpened ends are pressed together for 6 to 10 hours.

Belting Cement

Hide Glue	21/4 lb.
Water	$2\frac{1}{4}$ lb.
Glycerin	9 oz.
Carbolic Acid	3/ ₁₆ oz.

To use, melt and apply hot to the leather belt and place the joint under pressure until the glue is thoroughly set.

Canvas Awning Cement U. S. Patent 2,011,218

Rubber Latex	10	oz.
Varnish	1	oz.
Citronella Oil	1/100	oz.
Nigrosine B Solution	1/100	

Textile Glue

(for Doubling of Cloth, Shirting, Drill)

,	6,
	15 oz.
	5-10 oz.
	2 oz.
	75 oz.
	*

Warm and stir together.

Jute or Burlap Sheet Binder British Patent 412,498

Gilsonite	11 lb.
Asphalt, Petroleum	23 lb.
Naphtha, Petroleum	35 lb.
Mineral Silicate Filler	15 lb.
Asbestos, Fibrous	15 lb.
Linseed Oil	2 lb.

Upholsterer's Paste

Prepare a

a. Calcium Chloride Solution (25° Bé.)

cleared by pouring off solution from settled dirt, and add 160 kg.

b. Potato-Starch 100 kg. Water 100 l.

(Heated to 60-65° C.)
This glue has a good binding power, but dries very slowly and is hygroscopic.

Fine Bookbinder's Paste

Dissolve in

Water, Boiling Trisodium Phosphate	100	1.
Trisodium Phosphate	15	kg.
Borax	2.5	kg.
or (Alum	10	kg.
and add with stirring, a	solution	of:
Water, Cold	120	1.
Starch	50	kg.
Warm until fluid.		

Upholsterer's and Bookbinder's Paste

a. Potato-Starch	50 kg.
Water, Cold	140 l.
b. Caustic Potash (50° Bé.)	6 kg.
Sodium Silicate	15 kg.
Water Cold	50.1

c. Acid to neutralize to weak alkalinityd. Rosin Soap, Warm Fluid 5 kg.

Stir a till smooth, warm and stir with b to form a mucilage. Stir $\frac{3}{4}$ to 1 hour more, add c, then d, and stir slowly.

Bookbinder's Paste

a.	Rye or	Wheat Flour	100 kg.
	Water.	25° C.	200 1.

b. Caustic Soda (35° Bé.) 24 kg.
c. Nitric Acid until neutral

c. Nitric Acid
d. Alum, Cold Saturated

Solution 20 kg. Stir a to dispersion, treat mildly with b, neutralize with c, and add d.

Adhesive Paste for Rubber-Cloth on Cardboard

a.	Gutta Percha, Finely Cut Carbon Disulphide	$\frac{18}{20}$		
	Benzene	10	g.	
	Turpentine Oil	10	g.	
b.	Colophony	42		

a is mixed and soaked several days, then add b with gentle warming.

Mending China, Pottery and Casts

Save all the pieces of the broken article and store where the edges will keep clean until the repair is made. If the edges become soiled they should be washed clean and allowed to dry. The edges may be sanded lightly if necessary to remove the soil. The worker should know where each piece belongs before the work is begun. Small pieces should be cemented together previous to the main repair. A sand box is convenient to hold pieces upright while making the repair leaving both hands free for the work. It is made by putting 8 inches of clean sand in a convenient sized box.

Have at hand the cement, rubber bands, a bowl of warm water, tissue and soft rags. One rag should be reserved for wiping the fingers. Do not work with sticky fingers. Be accurate. If some part is not true after having been put together, soak until the cement is dissolved, wash the edges and begin over. Warm water will dissolve plaster or whiting cement and turpentine or alcohol will dissolve others.

The most durable cement is pure white lead ground in linseed oil, so thick that it will barely spread smooth with a knife. After drying thoroughly (about three months) it makes a seam which is practically indestructible but the mend is very

conspicuous.

A less conspicuous cement is made of beaten egg white and sifted whiting or plaster of Paris. A small amount should be mixed at a time as it hardens quickly. In some cases it is just as satisfactory to brush the edges with beaten egg white and dust well with sifted plaster tied loosely in double mosquito netting. The pieces should be fitted together at once and held in place by rubber bands (placed lengthwise, crosswise and diagonally) wrapped loosely in tissue paper and buried in a sand box. Care should be taken that the break lies so that the weight of the sand will hold it together. Leave it in the box at least 24 hours. After a week the superfluous plaster may be scraped away.

Sometimes the rubber bands will not hold the pieces true on a stemmed article, a vase or a jug. In this case string six bands of the same size and strength upon a piece of tape. Tie the tape around the neck or base of the article before beginning the gluing. After the parts are joined slip another tape through the bands and tie above the fracture. The bands pulling in unison will hold the break together. The pressure on all mended fractures should be great enough to force out the tiny air bubbles which otherwise reflect light making the seam

conspicuous.

Universal Putty for Wood, Stone, Glass, Porcelain

(Dries after 24-30 hours)

a. Alabaster Gypsum
Gum Arabic
b. Cold Borax Solution, Saturated.

Stir until pasty.

Preserve Jar Sealing Wax Washes off easily with hot water. Paraffin Wax 35 g. Trihydroxyethylamine Stearate 3 g.

Paraffin Bottle Cap Adhesive U. S. Patent 1,964,380

Chicle	1	oz.
Dammar	1	02.
Petrolatum, Liquid	1/2	OZ.
Warm and stir until homo	geneor	as.

Bottle-Cap Varnish

Dissolve 2 oz. of red Sealing-wax in 5 oz. of denatured alcohol.

Seal for Bottles

Beeswax		5	g.
Carnauba	Wax	1	g.
Paraffin		1	g.
Minium		5	ġ.
Whiting		2	ġ.

To Seal Glass Tubing to Iron Tubing Grind the ends you wish to join to a tapered fit and then seal by fusing with silver chloride.

Cement for Vacuum Tubes

Marble Flour	85	oz.
Shellac	10	oz.
Rosin	5	oz.
Phenol Formaldehyde Resin	25	oz.

Glass to Metal Seals

Formula No. 1

Iron	37	oz.
Nickel	30	oz.
Cobalt	25	oz.
Chromium	8	oz.

The above is suitable for use with lead-

No. 2

Iron	54 lb.
Nickel	28 lb.
Cobalt	18 lb.

Suitable for use with Corning glasses.

Safety Glass Adhesive U. S. Patent 2,009,029

Formula No. 1

A small portion of casein is heated in an open vessel with twice its weight of glycerol and 1.0% by weight sodium hydroxide (based on the casein). The temperature is brought gradually to 150-165° C. over a period of 15 minutes with continual stirring, and then held at this point for an additional 30 minutes. This product is a clear liquid at 100° but rubbery and very slightly opaque on cooling to room temperature. This material while hot may be pressed between two hot pieces of glass until air bubbles disappear, On cooling a piece of sandwich

glass is obtained in which the glass plates are firmly held together.

No. 2

Fourteen and nine-tenths (14.9) parts glycerol, 35.1 parts phthalic anhydride and 10.0 grams sheet gelatin (broken into small pieces) are heated with stirring in an open aluminum vessel, one hour up to 200° C. and 4 hours at 200° C., or to an acid number of 65–70. Some difficulty may be experienced in the early stages in making the bulky masses of gelatin mix with the other materials. This resin may be used as the sandwiching material for glass, or dissolved in a solvent such as acetone and used as an adhesive or impregnating agent.

Percent Quartz					
Coefficient of Expansion					
Percent Porcelain					
Coefficient of Expansion					

The quartz cement mixtures for values of quartz between 40% to 70% usually shows the same coefficient of expansion as pure cement. The modulus of elasticity of the quartz cement mixture increases with increasing quartz content. The bending strength, however, decreases almost in proportion to the percent quartz. The impact or shock bending strength, however, is practically unaffected up to 50% quartz content.

Porcelain and metal surfaces should be given a coating of a good elastic varnish before cementing. The cement should be allowed to harden in a steam chamber or, at least, be kept thoroughly wet for the

first forty-eight hours.

Another good porcelain cement is the usual litharge glycerin cement. This should be made in a ratio of three parts litharge and 1 part glycerin by weight. The glycerin used should contain less than 15% water and the litharge must, as far as possible, be free of lead carbonates as they produce a porous, weak cement.

A filler of up to 40% crushed or powdered porcelain may alse be used advantageously with the litharge. All exposed surfaces of cement should be given a thoroughly protecting coating of a good grade of Glyptal or Bakelite varnish.

1.

Litharge and glycerin ratio about 75/25 sample poured in a 25 mm. diameter glass tube hardens to a solid mass in less than 24 hours, but on further drying gives off additional moisture thereby slightly decreasing its dimensions so that it can be pushed out of tube. Swells

Mastic Seal for Oil Drums German Patent 613,748

Aluminum Powder	30 kg.
Nitrocellulose	14 kg.
Butyl Acetate	21 kg.
Ether	35 kg

Glass Electrical Cements

To offset the greater thermal coefficient of expansion of ordinary cement (11.5×10^{-6}) against that of porcelain (4.5×10^{-6}) a mixture of cement and powdered quartz or cement and crushed porcelain may be used. The thermal coefficient of expansion has approximately the following values:

0	20	40	70	80
11.5	10	8.5	5.5	4×10^{-6}
0	20	40	60	80
11.5	10.5	9	7.5	6×10^{-6}

on moist days sufficiently to firmly hold sample in glass tube. It is now adhering to glass. Under the microscope it shows a fairly dense even mass with numerous minute air-bubbles which appear to be coated with a shiny scale. Cracks when heat is locally applied and apparent traces of glycerin start to burn with a slow glowing, causing bubbles to be formed. Mechanically very rigid and strong, water absorption in 14 hours—1.6% by weight.

2

Equal parts litharge and crushed porcelain plus glycerin to make a good flowing cement. Hardens in less than 24 hours, forms a hard solid body which cannot be moved in glass tube but under the microscope shows somewhat more porous than No. 1, especially around the coarser grains of crushed porcelain. Mechanically rigid and strong.

3.

Glens Falls Cement Company iron clad portland cement and water. Cement poured in 25 mm. diameter glass tubes, hardens in less than 24 hours but 7 days is recommended by the manufacturer to give it full strength. One test tube was kept under water for the first 48 hours according to the recommendation of the manufacturer and one tube air dried The air dried cement could be hammered out of glass tube and under the microscope showed minute air bubbles imbedded in the solid material. sample set under water showed a very dense homogeneous body composed of minute bright crystals imbedded in a

mass of various dull colored material. The sample set under water showed considerable more strength and toughness than the air dried absorption in 14 hours —8.8% by weight.

4.

50% "iron clad" portland cement, 50% crushed porcelain. Sufficient water to readily pour sample set under water for the first 48 hours and allowed 6 days for air hardening. This sample gave a hard tough body of high mechanical strength. Under the microscope it showed the porcelain particles very densely imbedded in the material and traces of air bubbles could only be found around the larger porcelain grains. It appears to be a very promising cement for porcelain cementing. Number 4 very closely resembles the so-called "Teleo" Cement patented by the porcelain factory Treiberg in Thyringen, Germany, and consisting of portland cement and crushed quartz glass. This cement was developed with a view of obtaining a cement of approximately the same temperature expansion as that of porcelain. This is obtained by mixing a sufficient quantity of crushed quartz glass with an expansion coefficient of 0.5×10^{-6} with the portland cement having an expansion of 11.5×10^{-6} to give an expansion coefficient of approximately equal to that of porcelain of 4.5 × 15-6. Further tests on the va- 4.5×15^{-6} . rious cements are necessary in order to fully determine the mechanical properties.

Summary

The indications from the above preliminary tests, therefore, are that litharge and glycerin in a ratio of about 80/20 by weight or a mixture of 7 parts Glens Falls iron clad cement and 3 parts powdered porcelain or perhaps still better powdered quartz and water is the most suitable cement to use for bushing work.

The metal and porcelain surfaces to be given one coat of clear "Valspar" varnish to take care of the variation in expansion and all free surfaces of the cement to be given two or three coats of varnish as a protection against moisture.

To Plug Holes in Metal

Mix powdered sulphur and powdered aluminum 1-1 and pour on the metal which should be hot and clean. Then heat to melt the sulphur.

Metal Glue (for Tins, Etc.)

Resin (Shellac, Sandarac)

darac)

Manila-Kopal, Soft

50-100 g.

Manila-Kopal, Soft

Galipot or Turpentine,
Thick 3 g.
Alcohol, Denatured 100–200 g.
Castor Oil 1 g.

Pipe Joint Lute German Patent 597,044

Tallow 1 lb.

Mineral Oil 1 lb.

Melt together and mix with:

Ochre 1 lb.

Dry Clay or Sand 7 lb.

Premolded Expansion Joint
Chinawood Oil, Polymerized 5 lb.
Bitumen 85 lb.
Mineral Filler 10 lb.

Sulphur Thiokol Cements Formula No. 1

 Sulphur
 58.8 lb.

 Thiokol
 1.2 lb.

 Sand
 40.0 lb.

 No. 2
 Sulphur

 Thiokol
 1.2 lb.

 Sand
 38.0 lb.

 Carbon Black
 2.0 lb.

Refractory Cement U. S. Patent 1,952,119

Magnesium Oxide, Powdered
(Deadburned) 50 lb.
(Fused) 15 lb.

Zircon Sand
60-mesh 25 lb.
300-mesh 10 lb.

Sodium Silicate (d. 1.3) sufficient to make paste.

High Temperature Luting Compound
Alumina 50 lb.
Magnesia 25 lb.
Kaolin 25 lb.
Sodium Silicate sufficient to bring to a
working consistency.

Nitric Acid Resistant Putty

White Asbestos Powder
Blue Asbestos Fiber
China Clay
Linseed Oil

20 parts
10 parts
21 parts
22 parts

A cement for nitric acid plants contains:

Blue Asbestos Powder, and Sodium Silicate 1.5 Tw. Asbestos Binder U. S. Patent 2,010,224

Shellac 48 oz. Dicyandiamide 2 oz.

Heat together and stir until uniform.

Acid-Proof Dental Cement

Make a concentrated solution of silicate of soda and form a paste with powdered glass. Invaluable where a luting is required to resist the action of acid fumes.

Dental Cement British Patent 430,624

	,
Lithium Phosphate	½ oz.
Phosphoric Acid	5 oz.
Zinc Phosphate	$\frac{1}{2}$ oz.
Aluminum Phosphate	$\frac{1}{3}$ oz.
The above is added to a	ground porce-

lain of following composition:

A		
Alumina	30-50	oz.
Feldspar	10-20	OZ.
Sand	25-40	oz.
Zinc Oxide	1-10	oz.

Boiler Lagging

A splendid boiler lagging can be made by the following formula and applied direct to the boiler with a trowel, or molded into sections or blocks of suitable size and then dried and applied in the form of the usual sectional lagging:

1. 200 lb. spent Carbide Residue, drained to a soft putty consist-

2. 100 lb. Asbestos Fiber or Asbestos Fiber and Magnesia. (Old lagging properly ground will be satisfactory.)

3. 50 lb. Fine Dry Pine Sawdust.

Mix 2 and 3, then add 1 and mix thoroughly. If too dry add a small quantity of water. If oak or wet sawdust is used, quantity should be increased in the same proportion as the difference in weight per cubic foot.

It has also been found that carbide residue mixed with equal parts of Fuller's Earth will produce a good heat insulator for small furnaces.

Silicate Cements

Composition

Silicate of Soda and As-

Silicate of Soda and Sil-

Silicate of Soda and

Silicate of Soda and Dia-

tomaceous Earth

Silicate of Soda

bestos Fiber

ica or Clay

Whiting

Methods

Apply to porous surface and wash with dilute sulphuric acid after

setting

Mix to paste and wash with dilute sulphuric acid to develop acidresistance after setting

Mix to paste and wash with dilute sulphuric acid to develop acid-resistance after setting

Mix to paste and wash with dilute sulphuric acid to develop acid-resistance after setting

Mix to paste and wash with dilute sulphuric acid to develop acid-resistance after setting

Very quick setting; make only as needed

Portland cement may be added

Remarks

Acid-proofing of wood, unglazed tile, etc.

General acid-proof cement and lute; also used for setting acid-

proof bricks, etc.

Acid-proof and refractory

Silicate of Soda and Portland Cement

Silicate of Soda and Zinc Oxide, with or without added Clay

Silicate of Soda and Sawdust or Wood Flour

Silicate of Soda and Copper Powder For setting acid-proof tiles; waterproof

Used as a binder in abrasive wheels; water-resistant

Strong bond; water-resistant; also resistant to weak acid

For protecting spots during case hardening

*	Silicate Cements-Continued	l
Composition	Methods	Remarks
Silicate of Soda and Barytes Flour	Make to a stiff paste	Resists wet chlorine
Silicate of Soda and Duriron Dust	Make to a stiff paste	Used for temporary repairs of Duriron
Silicate of Soda and Silica Flour and Sodium Fluosilicate	Make to a stiff paste	Used for temporary repairs of Duriron
Silicate of Soda and 20 Manganese Dioxide; 20 Zinc Oxide; 10 Kiesel- guhr; 3 Graphite	Make to a stiff paste	Used for repair of metal parts; becomes highly acid resistant on setting.
	Glycerol-Litharge Cements	
Composition	Methods	
a. Glycerol and Litharge		Remarks
	Mix to a paste and apply promptly; varying the proportions, changes characteristics	Proportions vary; addition of water to glycerol hastens setting (2 water to 5 glycerol
b. (a) plus Whiting	Slower setting than straight cement	sets in 10 minutes)
c. (a) plus Silica	Slower setting than straight cement	
d. (a) plus Ferric Oxide	Slower setting than straight cement	
e. 1 part Litharge; 1 part Silica; 1 part Portland Cement, Gly- cerol and Silicate of Soda (diluted)	Addition of silicate con- trols sefting time	Sulphite digester linings; dilute sulphuric acid
f. 1 part Litharge; 1.5 parts Silica; 1.5 parts Portland Cement; Gly- cerol and more Silicate	Mix to a putty consist- ency	For sulphur dioxide gas (wet); resists hot solutions
g. 5 parts Litharge; 3 parts Silica; 2 parts Quartz Flour; c.p. Glycerol	Mix to a putty consist- ency	For sulphur dioxide gas (wet); resists hot solutions
 h. Glycerol and Litharge plus Graphite 	Mix to a putty consist- ency	Used on pipe joints which can be taken
i. Glycerol and Red Lead	Mix to a putty consist- ency	apart easily Acid-resistant joints in iron; sets hard
	Miscellaneous Cements	
Composition	Methods	70 -
Iron Filings (100); Ammonium Chloride (1); water	Mix to a thick paste	Remarks Used to repair cast iron, etc.; resistant to heat but not acids
Asbestos Wicking and Rubber Cement (rub- ber dissolved in ben- zene)	Soak wicking in cement and force into joint (not too strongly)	Used as caulking on fused silica and stone-ware bell and spigot joints; proof against moisture and dilute acids; flexible

Miscellaneous Cements-Continued

Composition	Methods	Remarks
Lead Wool	Caulk into joints	Used in the same way as poured lead joints in bell and spigot pipe
Asbestos Wicking	Used as a caulking with or without asphalt or other cement to protect it	Resists common acids ex- cept hydrofluoric
White Lead and Varnish Putty	.25 to 1.5 gal. of hard drying varnish to 100 lb. paste white lead in linseed oil	For jointing marble, stone, glass, etc.; an adhesive, slow-harden- ing cement
White Lead Paste with Read Lead	Red lead added to give the heaviest workable paste	For threaded pipe joints; can be opened
Lead Filings	Lead is filed on to pipe threads moistened with lubricating oil	Makes tight threaded joints
Red Lead in 3 parts, raw Linseed and 1 part medium Lubricating Oil	Mix to stiff paste	Adheres tenaciously to metal; remains soft and elastic; fillers may be added
Cellulose Acetate solu- tions (with or without fillers)	Applied as a sealing com- pound	General service adhesive
Cellulose Nitrate solu- tions (with or without fillers)	Applied as a sealing com- pound	General service adhesive
Rubber, Linseed Oil, Asbestos Fiber	Rubber is dissolved in hot oil and asbestos added to make a thick putty	For joints in stoneware, etc.; forms an elastic mass
Sulphur in various mix- tures	Sulphur is melted and mixed with clay, silica, etc., to form a putty	Applied hot as a grout- ing; resists acids and alkalis
Self-vulcanizing Rubber Cement	Painted or trowelled in place	Resists both corrosion and abrasion
Numerous resin base pro- prietaries		Resist dilute acids
Synthetic resin varnishes		Resist acids and weak al- kalies
Soaps (particularly of heavy metals)	Made to a putty with lin- seed or other drying oil	Resists hydrocarbon solvents
3 lb. dry White Lead; 2 lb. White Lead in Oil; 1 lb. 85% Magnesia with Linseed Oil to make stiff putty	Laid between flanges of joints, using a lead wire as a shim	Resists hot alcohol vapors
80 lb. Litharge; 8 lb. Red Lead; 10 lb. Floc Asbestos; 1.5 gal. Lin- seed Oil	Hardens in about 7 days	Resists dilute nitric acid cold but not hot
Tar or Soft Pitch and Linseed Oil (50-50)	Applied hot	Does not harden; resists acids
Sulphur melted with Rosin Tar or Pitch	Melted in place	Resists hydrochloric acid

Silicate of Soda and Barytes Flour Silicate of Soda and Barytes Flour Silicate of Soda and Duriron Dust Silicate of Soda and Silicate of Soda and Silicate of Soda and Silicate of Soda and 20 Manganese Dioxide; 20 Zinc Oxide; 10 Kieselguhr; 3 Graphite Composition a. Glycerol and Litharge Composition b. (a) plus Whiting C. (a) plus Silica d. (a) plus Ferric Oxide c. 1 part Litharge; 1 part Silica; 1 part Portland Cement, glycerol and Silicate of Soda (diluted) f. 1 part Litharge; 1 part Portland Cement, glycerol and Silicate of Soda (diluted) f. 1 part Litharge; 2 parts Court Soda (diluted) f. 1 part Litharge; 3 parts Silica; 2 parts Quartz Flour; c.p. Glycerol and Exilica; 2 parts Quartz Flour; c.p. Glycerol and Exilica; 2 parts Quartz Flour; c.p. Glycerol and Red Lead Mix to a putty consistency Mix to a thick paste Mix to a		Silicate Cements—Continued	
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A. Glycerol and Litharge a. Glycerol and Litharge b. (a) plus Whiting c. (a) plus Silica c. (a) plus Silica c. 1 part Litharge; 1 part Portland Cement, Glycerol and Silicate of Soda (diluted) f. 1 part Litharge; 1 part Portland Cement, Glycerol and Silicate of Soda (diluted) f. 1 part Litharge; 15 parts Portland Cement; Glycerol and more Silicate ency f. 5 parts Litharge; 3 parts Silica; 2 parts Quartz Flour; c.p. Glycerol h. Glycerol and Litharge plus Graphite i. Glycerol and Red Lead Mix to a putty consistency Mix		Glycerol-Litharge Cements	
Mix to a paste and apply promptly; varying the proportions, changes characteristics b. (a) plus Whiting c. (a) plus Silica d. (a) plus Ferric Oxide e. 1 part Litharge; 1 part Portland Cement, Glycerol and Silicate of Soda (diluted) f. 1 part Litharge; 1.5 parts Portland Cement, Glycerol and more Silicate ency g. 5 parts Litharge; 3 parts Silica; 2 parts Quartz Flour; c.p. Glycerol h. Glycerol and Litharge plus Graphite i. Glycerol and Red Lead Mix to a putty consistency Mix to a putty consis	Composition		Remarks
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c. 1 part Litharge; 1 part Silica; 1 part Portland Cement, Gly- cerol and Silicate of Soda (diluted) f. 1 part Litharge; 1.5 parts Silica; 1.5 parts Portland Cement; Gly- cerol and more Silicate g. 5 parts Litharge; 3 parts Silica; 2 parts Quartz Flour; c.p. Glycerol h. Glycerol and Litharge plus Graphite i. Glycerol and Red Lead Mix to a putty consist- ency Used on pipe joints which can be taken apart easily Acid-resistant joints in iron; sets hard Used to repair cast iron, etc.; resistant to heat but not acids Used as caulking on fused silica and stone- ware bell and spigot joints; proof against moisture and dilute	d. (a) plus Ferric Oxide		
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parts Silica; 2 parts Quartz Flour; c.p. Glycerol h. Glycerol and Litharge plus Graphite i. Glycerol and Red Lead Mix to a putty consistency Acid-resistant joints in iron; sets hard Used to repair cast iron, etc.; resistant to heat but not acids Used as caulking on fused silica and stoneware bell and spigot joints; proof against moisture and dilute	parts Silica; 1.5 parts Portland Cement; Gly-		(wet); resists hot so-
plus Graphite ency Mix to a putty consistency Noticlerance stant joints in iron; sets hard Used to repair cast iron, etc.; resistant to heat but not acids Used as caulking on fused silica and stoneware bell and spigot joints; proof against moisture and dilute	parts Silica; 2 parts Quartz Flour; c.p.		(wet); resists hot solu-
Mix to a putty consistency Notice repair cast iron, etc.; resistant to heat but not acids Used as caulking on fused silica and stoneware bell and spigot joints; proof against moisture and dilute			which can be taken
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zene) (not too strongly) ware bell and spigot joints; proof against moisture and dilute	monium Chloride (1); water		etc.; resistant to heat
	Rubber Cement (rubber dissolved in ben-	Soak wicking in cement and force into joint (not too strongly)	fused silica and stone- ware bell and spigot joints; proof against moisture and dilute

Miscellaneous Cements-Continued

Composition	Methods	Remarks
Lead Wool	Caulk into joints	Used in the same way as poured lead joints in bell and spigot pipe
Asbestos Wicking	Used as a caulking with or without asphalt or other cement to protect it	Resists common acids ex- cept hydrofluoric
White Lead and Varnish Putty	.25 to 1.5 gal. of hard drying varnish to 100 lb. paste white lead in linseed oil	For jointing marble, stone, glass, etc.; an adhesive, slow-harden- ing cement
White Lead Paste with Read Lead	Red lead added to give the heaviest workable paste	For threaded pipe joints; can be opened
Lead Filings	Lead is filed on to pipe threads moistened with lubricating oil	Makes tight threaded joints
Red Lead in 3 parts, raw Linseed and 1 part medium Lubricating Oil	Mix to stiff paste	Adheres tenaciously to metal; remains soft and elastic; fillers may be added
Cellulose Acetate solu- tions (with or without fillers)	Applied as a scaling compound	General service adhesive
Cellulose Nitrate solu- tions (with or without fillers)	Applied as a sealing com- pound	General service adhesive
Rubber, Linseed Oil, Asbestos Fiber	Rubber is dissolved in hot oil and asbestos added to make a thick putty	For joints in stoneware, etc.; forms an elastic mass
Sulphur in various mix- tures	Sulphur is melted and mixed with clay, silica, etc., to form a putty	Applied hot as a grout- ing; resists acids and alkalis
Self-vulcanizing Rubber Cement	Painted or trowelled in place	Resists both corrosion and abrasion
Numerous resin base pro- prietaries		Resist dilute acids
Synthetic resin varnishes		Resist acids and weak al- kalies
Soaps (particularly of heavy metals)	Made to a putty with lin- seed or other drying oil	Resists hydrocarbon solvents
3 lb. dry White Lead; 2 lb. White Lead in Oil; 1 lb. 85% Magnesia with Linseed Oil to make stiff putty	Laid between flanges of joints, using a lead wire as a shim	Resists hot alcohol vapors
80 lb. Litharge; 8 lb. Red Lead; 10 lb. Floc Asbestos; 1.5 gal. Lin- seed Oil	Hardens in about 7 days	Resists dilute nitric acid cold but not hot
Tar or Soft Pitch and Linseed Oil (50-50)	Applied hot	Does not harden; resists acids
Sulphur melted with Rosin Tar or Pitch	Melted in place	Resists hydrochloric acid

Miscellaneous Cements-Continued

Composition

Shellac (30); Rosin (20); Alcohol (33); Gypsum (2); Ferric Oxide (15)

2 parts Scotch Glue; 7 parts Water; 1 part Glycerol

Methods

Remarks

Resists petroleum oils

Finely powdered so	Trub
are mixed with an	
coholic solution of	the

For oil or gas leaks; more glycerol softens it

Non-Efflorescing Concrete
The addition of 5% Barium Carbonate
to the cement prevents efflorescence.

Keying Plaster to Concrete

First secure a fast setting plaster which corresponds to Plaster of Paris, moulding plaster or something similar. This plaster is mixed thin enough so it can be whipped onto the wall with a brush. After this dash coat of plaster has thoroughly set, the wall, which now has a rough surface, may be plastered over in the usual way with ordinary gypsum plaster.

Plaster Cement, Patching U. S. Patent 2,016,986

Calcium Carbonate (50–200		
mesh)	4 1	b.
Dry by heating below 600° C. Slaked Lime		
Slaked Lime	5 1	b.

Refrigerator Display Case Caulking Compound

U. S. Patent 1,974,745

Nitrocellulose	1-7	oz.
Dibutyl Phthalate	15-60	oz.
Asbestine (Mineral)	30-90	oz.
Camphor	1/4	oz.

Cement "Wash" Hardener
Portland Cement 20 lb.
Iron Filings 126 lb.
Water 9 lb.

Concrete Wash, or Finish Paint (Hard and Durable)

Apply with brush, mixing often.

•	,	
Slaked Lime		1 lb.
Cement		1 lb

Mix in water containing ½ lb. salt per gallon to desired consistency.

Colored Caulking Cement U. S. Patent 2,011,607

A cement of substantially permanent elasticity and which is adapted for ap-

plication by a trowel or a grease gun consists of paracoumarone resin m.p. about $50-60^\circ$ C. about 60, asbestos fiber about 20, a metallic oxide such as oxide of zinc or iron about 5 and xylol about 15%.

Pliable Glazing-Caulking Cement British Patent 398,057

Formula No. 1

Mineral Filler	1 - 50	oz.
Oil	30	oz.
Asbestos Fiber	20 - 1	oz.
Aluminum Powder	1-30	OZ.
Varnish sufficient to make	paste.	

No. 2

Calcium Carbonate, Powdered Magnesium Silicate,	12.60	oz.
Powdered	17.10	oz.
Asbestos Fiber	5.45	oz.
Soya Bean Oil	30.63	oz.
Varnish	16.22	02.
Aluminum Powder	9.00	oz.
Naphtha, Petroleum	9.00	oz.

Glazing Putty

- CAMBING	x acces		
Whiting, Domestic,			
200 mesh		205	lb.
Whiting, Belgian		70	lb.
Linseed Oil, Raw		26	lb.
Japan Drier		1	lb.
Mineral Spirits		3	lb.

Cement for Pestle Handles

Heat the head of the pestle until it is too hot to hold in the hand. Pour melted shellac into the hole, then take the wooden handle, wind some twine around the screw portion, and press it "home." Keep under pressure until the head of the pestle is cold.

Mortar Cement

Fuse together, in an iron vessel, equal parts of guttapercha and shellac. This forms a powerful cement. Strongly heat the edges of the broken mortar, apply a thin layer of the cement to both frac-

tured surfaces, and put together under pressure.

Joining Stainless Steel in Knife Handles Method 1

A waterproof cement is used, made by mixing finely powdered litharge and glycerin. The glycerin should be added in an amount equal in volume to half the volume of the powdered litharge and mixed thoroughly. The end of hollow handle is filled with cement and then insert the blade. Setting time about 45 minutes. Mix only enough cement as needed as it sets quickly becoming hard and insoluble.

Method 2

The stainless steel blade is first thoroughly tinned and then soldered in place. It is necessary to have all parts clean and free from scale. Solders used are either 50% tin and 50% lead or 66% tin and 34% lead. Flux used is made up of zinc chloride, commercial grade, 37 g.; glacial acetic acid 99.9%, 23 g.; hydrochloric acid (commercial), 34.5%, 40 g.

Metal Adhesive

	Nitrocellulose Scrap	10	g.
	Alcohol	26	g.
	Ethyl Acetate	25	g.
	Butyl Acetate	31	
	Benzol	30	
	Camphor	2	g.
	To the viscous solution add:		
	Metal Powder enough to "	hide	,,
	Viscosity should be high en		to
p	revent the metal settling down	۱.	

Rubber to Metal Cement British Patent 432,493

Paris White	40	oz.
Rosin	3	oz.
Dammar or Copal Gum	15	oz,
Benzol	15	oz.
Naphtha	23	OZ.
Rubber	11/2	oz.

Pyroxylin to Metal Adhesive

Pyroxylin	6 oz.
Gelatin	7 oz.
Acetic Acid, Glacial	87 oz.

Aluminum Foil to Leather or Paper Adhesive

U. S. Patent 1,925,903

Linse	d Oil	Fatty	Acids	11.82	g.
Tung	Oil	•		16.35	
Rosin				22.53	ğ.
Hont	monid	lr in	olumin.	**** ******	1 +

280° C.; cool to 260-265° C. and add with stirring:

Phthalic	Anhydrid	e	32.68	g.
Glycerin	_		16.35	g.
Ethylene	Glycol		4.22	
Troop of	0000 000	α	 olonn.	hoot

Keep at 200-220° C. until clear; heat at 250° C. until a sample solidifies in 40 seconds at 200° C.

Take of the above resin	11 g.
and dissolve in: Acetone	11 g.
Dibutyl Phthalate Nitrocellulose "Solution"	5 g.
(½ second)	sufficient

Thermoplastic Cement

TAZOTAKO PAGETO GALLONIO			
Nitrocellulose Wet 5-6 sec.	8	g.	
DuPont Resin RH-35			
6# cut	10	g.	
Dibutyl Phthalate	4	g.	
Methyl Ethyl Ketone	10	g.	
Butyl Acetate	10	g.	
Toluol	58	g-	

Fusible Adhesive Cement U. S. Patent 1,945,803

Chlorinated Naphthalene		
(Solid)	50	oz.
Ester Gum	50	0Z.
Rubber Latex	อั	OZ.

Shellac Scaling Composition

Shellac	50	oz.
Beechwood Creosote	5	oz.
Ammonia (28%)	. 1	OZ,
Terpineol	2	oz.

Adhesive Sealing Compound (Universum)

Mix hot beeswax and Venice turpentine 1 to 1. Proportions may be varied according to needs. Can be colored if desired. This is very good to temporarily attach glass to iron or wood.

"Syndetikon" (Universal Adhesive)

	, (c	
a.	Prepare Caustic Lime,	
	Freshly Burned	100 g.
	Water	50 g.

Let stand to cool: pour off layer of water. Use now:

Lime Hydrate (above) 15 g. Sugar Solution (25%) 240 g.

Heat to 75° C., let stand stirring through from time to time, pour off the clear solution, of which

> Lime-Sugar Solution 225 g. Bone Glue 60 g.

Rosin 22.53 g. are mixed to swell over night. Dissolve finally by warming up.

Acid Resisting Cement Fine Sand 2 lb. Short Fiber Asbestos 2 lb. Magnesia 1 lb. Sodium Silicate sufficient to make paste. Aquarium Cement 3 lb. Litharge 3 lb. Fine White Sand 3 lb. Plaster of Paris Mix thoroughly. Then add linseed oil sufficient to make paste, and a small amount of drier. Adhesive Foil U. S. Patent 1,955,075 Acidify defibrinated blood at 40° C. with 0.5% lactic acid; mix with 2.3% ammonium sulphate solution; keep at 40° C. for 1-3 hours; render slightly alkaline and mix with 8-12% glycerin and 5% alum or synthetic tannins.

Adhesive for Casein Plastics British Patent 411,058

Casein	1	part
Water	1	part
Urea	$\frac{1}{2}-1$	part

Quick Hardening Putties German Patent 613,748

Formula No. 1 Aluminum Powder g. Nitrocellulose 14 g. Butyl Acetate 21 cc. Ethvl Ether cc.

No. 2		
Aluminum Powder Ethyl Cellulose Benzol Ethyl Ether	30 14 33.6 22.4	
-		

Red Lead Putty

Red Lead, Dry	31 lb.
White Lead, Dry	48 lb.
Silica	16 lb.
Raw Linseed Oil	1 gal.
The state of the s	

Slate Color Putty		
Whiting	24	lb.
White Lead, in Oil	70	lb.
Lampblack, Dry	2	oz.
Raw Linseed Oil	6	lb.

White Putty

Whiting	77	lb.
White Lead, in Oil	9	lb.
Raw Linseed Oil	14	lb.

Black Plastic Putty

400	lb.
100	lb.
7	gal.
	gal.
70	gal.
	100 7 7

Directions:

Melt the two blacks to 550° F. and hold until in complete solution, then add both oils and heat to 575° F. Cool to 450° and reduce.

The black fish oil is a very dark crude and cheap oil, unfiltered and full of stearines.

For overglazing where the lights of glass overlap, a semi-liquid coating is made by mixing into the base vehicles while hot 1/4 lb. of long-fiber asbestos to each gallon.

For the plastic putty for cementing the glass to the frame, the following mixture is made in a regular pony chaser:

Base Vehicle (above)		gal.
Stove Distillate	$1\frac{1}{2}$	gal.
"Asbestine"	50	Ĭb.
Long-fiber Asbestos	5	lb.

This product is stiff and must be applied by knifing or with a small trowel.

In the east and south cement slabs called cementiles are quite commonly used in constructing factory roofs. The joints of these tiles are first partly filled in with a non-shrinkable cement, and above this flush with the tile surface is run a waterproof expansive plastic for protection. An eastern manufacturer of cementile roofing slabs also makes the joint cement or putty. They buy large quantities of paint skins from paint manufacturers, and use this as the base material, cooking the same with an addition of fish oil, subsequently churning it with such filling material as asbestine or whiting, short asbestos fiber, and red oxide for color. The final protective is a wellknown commodity, trade name similar to "mud mud." Its salient features are: a soft but firm plasticity; a condition of slime for easy slip in trowelling; slow setting during manipulation, but later becomes surface-set out of dust and dirt: retains its softness and cohesiveness within the joint, indefinitely. These features have been very well reproduced in the following formulation:

5% Leaded Zinc Oxide	24	lb.
Borate of Manganese	1/2	lb.
Spanish Red Oxide	8	lb.
Treated China Wood Oil	4	gal.
Sulphurized Fish Oil	4	gal.
Medium Body Gloss Oil	4	gal,
"C" Asbestos Fiber	32	lb.

The prepared oil is 40 lb. of limed rosin and 20 gal. of wood oil heated to 425° F. and held there about 2 hours until very heavy—but no stringing; then reduced immediately with 50 gal. kerosene.

The above plastic is run into the tile joints with a hand-pressure caulking and glazing gun, fitted with either the standard or the extra large caulking nozzle.

Although akin to putty but more proporly termed otherwise, is that compound familiarly known by almost everyone as Litharge-Glycerin Cement, which is valuable for a number of purposes for which ordinary cement and putty would be neither practicable nor desirable. Probably all readers may feel that they know how to mix this cement for usage, but those who merely combine these two ingredients really would not be doing it efficiently for best results. The cement is correctly produced by adding to a mixture of 5 parts of 95% pure glycerin and 3 parts of water, sufficient finely ground litharge to form a plastic of any required consistency. Variation in the amount of water will influence the time of setting and to an extent the general characteristics, but all modification within the range of say 1 to 3 parts of water with 5 or 6 parts of glycerin will attain satisfactory hardness. Its normal hardening time is about ten minutes, but it may be made to remain soft for a longer period by an addition of ten per cent of inert material such as silica, iron oxide, or fuller's earth. Such admixtures do not detract from the ultimate hardening or strength, but also are beneficial in preventing possible cracking. Litharge-Glycerin Cement will withstand a high degree of combined heat and moisture. A very common usage is for forming water-tight connections between iron pipes and porcelain fittings; and for cementing glass aquariums, etc. Its most conspicuous feature is its resistance to practically all acids not of full strength. It is used to good advantage in temporarily sealing leaks at seams, around the bottoms, and around flanges, etc., of storage tanks filled with varnish; these temporary repairs have held until the contents of the tanks were used when a permanent repair could be made.

Marine Putty, to harden under water, may be made from the formulation here given:

Hydraulic Cement	30	lb.
Plaster of Paris	71/2	lb.
Litharge	10	lb.
Belgium Whiting	20	lb.

Lead Carbonate (Dry) 10 lh. Boiled Linseed Oil 3 gal.

On the seaboard, hydraulic cement is better known as sea-water cement. This type differs from regular Portland cement for land construction in being darker color and containing a minimum of tri-calcium aluminate... the constituent in cement which is rapidly attacked by (saline) sea water. Whereas regular cement contains 10-15% tri-calcium aluminate, this is minimized to 2% in seawater cement.

Painters' Lead Putties, also termed Hard Putty and Carriage Putty, will vary in lead content from almost straight lead to approximately 75 per cent and 50 per cent; the admixtures being whiting and/or silica. Typifying the first two, are the formulas below of hard putties actually used in railroad shops:

Dry White Lead	90	1b.	50	lb.
White Lead in Oil			20	lb.
Whiting			25	lb.
Silex	- 6	1b.		
Boiled Linseed Oil	3/16	gal.	-	
Gold Size Japan	3/16	gal.	3/4	gal.
Rubbing Varnish	$1\frac{1}{8}$	gal.	3/4	gal.

These mixtures are allowed to stand 72 hours to thoroughly wet down and sweat, and then kneaded up into putty. The silex used is the live quartz silica mainly adopted for the making of paste wood fillers. The pigmentation of a representative painters? hard putty with lower lead content would be 50% dry white lead, 35% whiting and 15% silica.

A non-shrinkable type of putty containing about 20% of lead in the pigment is this:

Whiting	125	lb.
White Lead, Dry	371/2	lb.
Silica	121/2	lb.
Raw Linseed Oil	31/2	gal.
Flour Paste	3½ 10¼	lb.

The flour paste is 2 lb. of wheat flour beaten up in about 1 quart of cold water and then poured into 3 quarts of boiling water, and boiled 5 minutes. Yield 1014 lb. net.

The foregoing non-shrinkable putty is very similar to what used to be known as Swedish putty, purported to be so excellent for wood, iron, or stone. Another type of Swedish Putty without lead, is the following:

2	1b.
	gal.
into	
1	gal.
then	stir

Whiting.	20	lb.
Whiting Gold Size Japan Raw Linseed Oil Grind in a paint mill.	50 2 1	lb. gal. gal.

Combine the two parts in a pony chaser, and thicken with more whiting to the required plasticity for knifing. This batch produces 100 lb. net.

Metal Furniture Baking Putty

THEOREM I WILL	our C I	wiring .	Lacoy
Bolted Whiting	:		5 lb.
mixed with Boiled Linseed then	Oil		1 pt.
Flour Paste			1 pt.

Mix all very thoroughly. The flour paste is as given for non-shrinkable putty. In all cases of preparing flour pastes, the flour and cold water should be beaten until entirely free from lumpiness; and during the subsequent cooking, should be continually stirred.

Stopping Putty is a dry mixture of 2 lb. of "Alabastine," 1 lb. of wheat flour, and 1 lb. of Portland Cement. When ready to use, 1 pound of this mixed powder should be thoroughly worked up to a stiff putty with 8 fluid ounces (1/2 pint) of cold water. This putty sticks to stone, wood, brick, etc.; used for filling knot holes, cracks, etc. Keep the dry powder in an air-tight jar.

Gesso Duro is Italian hard plaster used in making bas-relief casts. When dried, it becomes very hard and durable.

This product, per formula, below, remains soft and manipulable for quite a period of time, using a small trowel, spatula or by forming with the hands:

LePage's Fish Glue	4	gal.
Water, to reduce it	1	gal.
Oil of Lavender	6 fl.	. oz.
Raw Linseed Oil	1	gal.
Bolted Danish Whiting	50	lb.
Rubbing Varnish	1	gal.
Bolted Danish Whiting	20	lb.
(colors in oil may be adde ing is desired)	ed, if	shad-
ing is desired)		

Plastic Wood Dough

*Gum Solution		1 gal.
Glycerin		3 pt.
Butyl Alcohol		3 pt.
Whiting		8 lb.
Wood Flour		24 lb.
Dope (Solution)		8 gal.
Allera Co.		

"The "gum" solution is 16 pounds of gum rosin (WW Rosin) cold-cut (dissolved) in 1 gallon of methyl acetone; the "dope" is another cold-cut solution,

basis of 1 pound of "movie",-film scrap to each gallon of methyl acetone. The picture film scrap should be desilvered by washing in hot water to remove its gelatin coating and then laid out in the sun and air to dry; but preferably it is obtainable cleaned and ready for cut-

Onyx Cement

The above wood dough product is a soft workable putty easily applied to all kinds of depressions to be surfaced up. The work or job should not be left in too-rough state because the putty dries and hardens very rapidly; the ultimate sanding down later is a rather tough job unless the puttying had been reasonably smoothly applied.

There is one putty specially used in fair quantity, which is very little known in regular paint circles. This is termed Onyx Cement because its specific utility is for bonding slabs of onyx, marble, glass, and their imitations, to the walls in public buildings. It is necessarily of rather firm plasticity because of the weight it must partially support. Uniform handfuls of the putty are attached to the wall foundation at intervals about 18 to 24 inches apart; the slabs mentioned are then stood upright on their base, and then pressed back steadily and firmly into the mounds of putty. Suction, and the adhesive strength of the putty, securely hold the marble and glass permanently in place. The same material, plain or colored, is embedded in the joints between the slabs. The composition of this putty follows:

Domestic Whiting, 350 Mesh 100 lb. Domestic Whiting, 200 Mesh 100 lb. "Super-Sublimed" White 40 Lead lb. 1¼ gal. White Oil Drier Bodied Linseed Oil 11/4 gal. Boiled Linseed Oil 21/2 gal.

For certain work a Black Onyx Cement is used. This is produced on a bituminous base.

Another specialty probably even less known than the onyx putties . . . in paint circles, is a Black Packing Compound required by makers of corrugated iron culverts. These culverts are sturdy Armco-iron corrugated pipe, galvanized, in sizes from 12 to 84 inches diameter. They are the aqueducts for streams crossing the highways and for surface-sewers under driveways in rural districts, etc. There is first applied hot a thoroughlytested bituminous mastic pavement along the line of flow where erosion is greatest ... approximately the lower one-quarter or one-third of the inside circumference. This coating practically fills the valleys of the corrugations and to the extent of building up a thickness of perhaps ¼-inch over the rises.

For this purpose the culvert manufacturer supplies a plastic for cold application. The composition is 3 parts by weight of sawdust and 1 part asbestos fiber, thoroughly churned together with enough coal tar solution to form a putty that may be applied by hand to the abraded spots in the paved section of the

The last unusual specialty to be mentioned is Sheet Metal Deadener. Two eastern manufacturers have been supplying during the past three or four years a plastic compound developed for sounddeadening sheet metal equipment, principally metal furniture and automobile parts. This became most essential with the advent of the closed body, to eliminate rumble and vibratory noises, and especially the "tinny" sound caused by closing the doors. It is a standard application on Ford, Auburn, Stutz, Marmon, Duesenberg, and Nash cars; and probably on many others. The material might be described as a very soft bituminous plastic apparently containing fine asbestos fiber or other filler; it surfaces dust free very quickly, has excellent adhesion and undoubtedly maintains flexibility indefinitely. As general practice, it is applied onto the inner surfaces of the auto body and doors, or other object, to a thickness of approximately ¼-inch, using a trowel, broad knife, or spatula. This sets in less than 30 minutes, but soft; is firm in 1½ hours and still somewhat soft, is solid in 4 hours but not hard; and shrinks down somewhat in solidifying. For large production as by body builders and in the auto plants, the material has sufficient "slip" so it can be sprayed with special equipment.

High grade cork paint films insulate surfaces against heat, cold, and moisture, also deaden sound and soften the effect of shocks and blows, rendering them valuable for use on automobiles, railroad cars, and acroplanes. In the automobile industry they are employed to advantage on the lower sides of the engine bonnets and mud guards. Applied to the bonnets, they protect the outside lacquer films against the radiating heat of the motor; while the cork paint films on the lower sides of the mud guards protect the latter against the impact of stones, sand, etc. Applied to the surfaces of aeroplane cabins, they form a rather effectual insulation against the noise of the motors.

COATINGS, PROTECTIVE AND DECORATIVE

Marine Paints

Marine paints differ from house paints chiefly in that harder pigments are required. This means that such pigments as zinc oxide and iron oxide are used more extensively in marine paints than in house paints. Since steel vessels have largely replaced wooden vessels in seagoing traffic, the formulas shown herein are for the preservation and beautification of steel rather than wood. On steel the priming coat of paint—that is, the paint applied first on the metal-is of more importance than the priming coat on wood. The service to which marine paints are exposed is much more severe than that to which house paints are exposed. To meet this condition the various parts of the vessel must be considered separately. The paints suitable for the parts seen from the outside when the vessel is afloat are quite different from the paints suitable for underwater portions of the vessel. The paints suitable for inboard bulkheads are quite different from those suitable for inner bottoms or bilges, etc.

An excellent priming paint for steel surfaces to be exposed to the atmospheric elements is made from the following formula which produces one gallon and spreads approximately 650 sq. ft. per

gallon:	
Red Lead, Dry	20 lb.
Raw Linseed Oil	5 pints
Petroleum Spirits	2 gills
Paint Dryer	2 gills

Paint from the above formula should be used within a month after it is mixed. If allowed to stand in closed (or open) containers for an appreciably longer period, the pigment settles hard and cannot be again stirred to proper consistency for painting. By using very finely ground red lead pigment which contains 99 per cent true red lead, it is possible to successfully store the paint through periods of approximately one year. However, if the paint is to be stored during such period, or longer, formulas such as the following should be used:

Red Lead, Dry	1 lb.	11	oz.
Zinc Oxide, Dry		13	OZ.
Venetian Red. Dry	4 lb.	2	OZ.

Magnesium Silicate, Dry Spar Varnish 2 lb.	10	oz.
	7	oz.
Petroleum Spirits	9	oz.
Paint Drier	14	oz.
Aluminum Stearate	1	oz.

Films from paints of the above formulas interfere with the adhesion of shipbottom paints, so these paints should not be used on the outside underwater portion of the hull. If it is desired to prevent corrosion on that portion of the vessel during construction, a weaker film paint should be used, such as:

Metallic Brown, in Oil 7.5	lb.
Raw Linseed Oil 2.3	lb.
Spar Varnish .3	lb.
Gasoline .6	lb.
O?*	

- 01	
Metallic Brown, Dry	4.0 lb.
Spar Varnish	4.4 lb.
Paint Drier	2.5 lb.

The above two formulas are also suitable for a paint to be used on freshly pickled steel to protect it during fabrication; that is, as shop coat or field coat paints.

Aluminum paint may be used in lieu of red lead paint, for priming steel, but should not be used on underwater portions of the vessel. Its bright luster aids inspection of the interior of vessels under construction, but in warm, humid climates it does not prevent rust as does red lead paint. The formula is

_		Localina	~ ~~~		****		
		num Po				2	lb.
	Alumi	num M	ixing	Varn	ish	1	gal.
	Note:	This pa	aint s	hould	be	used	withir

a few hours after mixing.

While priming paints will give fair protection when used alone, they are designed to be covered with at least two coats of finishing paint. Unlike house paints, there is no advantage in using a different formula for the first and the second coat of marine finishing paint. Following are formulas for ten gallons of finishing paints—on surfaces not to be exposed underwater:

Outside White Paint

Titar	nox B,	in	Oil	3.0	85	lb.
Zinc	Oxide,	in	Oil		36	lb.

Ultramarine Blue, in Oil Raw Linseed Oil Petroleum Spirits Paint Drier	.5 oz. 30 lb. 3 lb.	Paint Drier Ultramarine Blue, in Oil	4 lb. .5 oz.
or White Lead, in Oil Zinc Oxide, in Oil Raw Linseed Oil Petroleum Spirits Paint Drier Ultramarine Blue, in Oil		Inside White Enar Titanox B, Dry Zinc Oxide, Dry Damar Varnish Pinc Oil Ultramarine Blue, in Oil	25 lb. 25 lb. 68 lb. 6 lb. .5 oz.
Oltramarine Blue, in Oli	1 oz.	To this white enamel me color pigments, ground in o	il or in var-
Outside Black Paint		nish, to produce desired shad	es. By add-
Lampblack, in Oil	88 lb.	ing additional pine oil just l	pefore apply-
Raw Linseed Oil	32 lb.	ing, the enamel is made	to brush on
Lampblack, in Oil Raw Linseed Oil Paint Drier	14 lb.	much easier. An enamel wil	not adhere
or		well over an enamel or gloss	y finish. If
Lampblack, Dry Spar Varnish Petroleum Spirits Paint Drier	4 lb.	two coats are to be applied, should be a flat paint.	me nrst coat
Spar Varnish	14 lb.	should be a nat paint.	
Petroleum Spirits	4 lb.		
Paint Drier	18 lb.	Outside Buff Pair	+
		TUDIA T and in O'l	10 TO T
Inside White Paint		White Lead, in Oil	125 lb.
Titanov R in Oil 5	6 lb.	Yellow Ochre, in Oil Venetian Red, in Oil Raw Linseed Oil Petroleum Spirits Paint Drier	14 lb.
Zinc Oxide, in Oil	1 ib.	Raw Lingged Oil	5 lb. 27 lb.
Raw Linseed Oil	4.5 lb	Petroleum Spirits	7 lb.
Zinc Oxide, in Oil Raw Linseed Oil Damar Varnish Petroleum Spirits Paint Drier	8 lb.	Paint Drier	4 lb.
Petroleum Spirits 2	0 lb.		1 10.
Paint Drier	4 lb.		
Paint Drier Ultramarine Blue, in Oil	.5 oz.	Inside Semi-flat Light Gre	en Paint
Or	1		
White Lead, in Oil 7	7 lb.	Titanox B, Dry Zinc Oxide, Dry Chrome Green Oxide, in Oi Damar Varnish	30 lb.
Zinc Oxide, in Oil 7	7 lb.	Chrome Green Oxide, in Oi	1 7 oz.
Raw Linseed Oil 1	8 lb.	Damar Varnish	39 lb.
Petroleum Spirits 1	5 lb.	Damar Varnish Petroleum Spirits	20 lb.
Paint Drier 3	4 lb.		
White Lead, in Oil 7 Zinc Oxide, in Oil 7 Raw Linseed Oil 1 Petroleum Spirits 1 Paint Drier 3 Ultramarine Blue, in Oil	.5 oz.	7 12 7	
	1	Inside French Gray Er	namel
Light Gray Paint		Titanox B, in Oil Lampblack, in Oil	72 lb.
Titanox B, in Oil 50 Zinc Oxide, in Oil 35 Lampblack, in Oil IUltramarine Blue, in Oil Raw Linseed Oil 39 Petroleum Spirits 1 Paint Drier 8	1b.	Lampblack, in Oil Chrome Yellow, in Oil Spar Varnish Damar Varnish Pine Oil	1 lb.
Zinc Oxide, in Oil 35	lb.	Chrome Yellow, in Oil	1 lb.
Lampblack, in Oil	. lb.	Spar Varnish	30 lb.
Ultramarine Blue, in Oil	34 lb.	Pine Oil	29 lb.
Raw Linseed Oil 39	lb.	I life Oil	6 lb.
Petroleum Spirits 1	.5 lb.		
Faint Drier 8	16.	Piping, ducts, gas cylin	iders, etc.,
		aboard vessels are usually m	arked with
Outside Green Paint		colors to indicate the purpos	e served or
Chrome Green, Dry 30	o Ib.	the contents. Formulas for are:	such paints
Zinc Oxide, Dry) lb.	Red Paint	
Chrome Yellow, in Oil	3.6 lb.		
Chrome Green, Dry Zinc Oxide, Dry Chrome Yellow, in Oil Yellow Ochre, Dry	7.5 lb.	Toluidine, Dry	7 lb.
Lampotack, in Oil	i 1b.	Spar Varnish	73 lb.
Spar Varnish			
Petroleum Spirits 10		Blue Paint	
Paint Drier	lb.		400
		White Lead, in Oil	106 lb.
Inside Flat White Paint	- 20	Ultramarine Blue, in Oil	26 lb.
Zinc Oxide, in Oil 157	7 lb.	Raw Linseed Oil Petroleum Spirits	22 lb.
Petroleum Spirits 23		Paint Drier	8 lb.
		- WARRY BUT ALVE	4 lb.

Green Paint Chrome Green, in Oil Raw Linseed Oil Petroleum Spirits Paint Drier	97 lb. 21 lb. 9 lb. 5 lb.
Black Paint Lampblack, in Oil Petroleum Spirits Paint Drier	70 lb. 9 lb. 10 lb.
Brown Paint Metallic Brown, in Oil Raw Linseed Oil Petroleum Spirits Paint Drier	100 lb. 27 lb. 8 lb. 3 lb.
Yellow Paint Chrome Yellow, in Oil Raw Linseed Oil Petroleum Spirits Paint Drier	116 lb. 20 lb. 9 lb. 3 lb:

The above red and green paints are suitable for the stands on which running lights are mounted, red marking the port side and green the starboard side.

Single shell smoke stacks become too hot for any of the above paints. Such surfaces should be painted with special paints, the following formulas being typical:

ypical:		
Light Gray Paint		
White Lead, Dry	48	lb.
Zinc Oxide, Dry	19	lb.
Litharge, Dry	3.5	lb.
Lampblack, in Oil	5	lb.
Ultramarine Blue, in Oil	.5	lb.
Damar Varnish	20	lb.
Kerosene	33	lb.
Paint Drier	6	lb.
or		
Titanox B, Dry	60	lb.
Interior Varnish	52	lb.
Lampblack, in Oil	2	lb.
Petroleum Spirits	9	lb.
(a) () () () () () () () () (
Red Paint		
Indian Red, Dry	40	lb.
Interior Varnish	55	lb.
Paint Drier	15	lb.
X		
Buff Paint		
White Lead, Dry	55	lb.
White Lead, in Oil	55	lb.

18 lb.

13 lb.

12

23 lb.

.5 lb.

lb.

lb.

Silica

Yellow Ochre, in Oil

Venetian Red, in Oil

Boiled Linseed Oil

Petroleum Spirits

Litharge, Dry

Green Paint			
Chrome Green, Dry	30	lb.	
Lampblack, in Oil	2	lb.	
Interior Varnish	60	lb.	
Paint Drier	10	lb.	
Black Paint			
Drop Black, Dry	38	lb.	
Interior Varnish	48	lb.	
Paint Drier	16	lh	

The waterline area on the outside of the hull is generally regarded as the most difficult part of the vessel to keep properly painted. This is because it is subjected to both atmospheric and underwater exposure, and paints suited to the one exposure are not suited to the other. A high grade varnish paint applied over red lead primer gives as good service on this area as has been obtained. Typical of waterline paints are:

Red Paint	
Venetian Red, Dry	29 lb.
Spar Varnish	42 lb.
Petroleum Spirits	7 lb.
Paint Drier	18 lb.
-	

The state of the s	
Light Gray Paint Zinc Oxide, Dry Lampblack, in Oil Ultramarine Blue, in Oil Spar Varnish Petroleum Spirits Paint Drier	30 lb. 8 lb. 12 lb. 31 lb. 17 lb. 18 lb.
Black Paint	
Drop Black, in Oil Zinc Oxide, Dry Spar Varnish Petroleum Spirits Paint Drier	20 lb. 19 lb. 20 lb. 20 lb. 17 lb.
Lampblack, Dry Zinc Oxide, in Oil Spar Varnish Petroleum Spirits Paint Drier	8 lb. 20 lb. 38 lb. 7 lb. 18 lb.

Shipbottom paints are used to prevent rust and to prevent the attachment of marine fouling on the bottoms of vessels. The "anti-corrosive" paint is to prevent rust and is applied next to the steel. The "anti-fouling" paint is to prevent the attachment of barnacles, algae, and other forms of fouling. It contains material toxic to marine organisms, and is applied over the anti-corrosive paint. Both paints should be quick drying paints. Each of the two paints is so dependent on the

other that the two formulas are shown
together. The anti-corrosive paint of one
set should not be used with the anti-
fouling paint of another set. The follow-
ing formulas are typical:

Anti-corrosive Paint

Gum Shellac	8 lb.
Denatured Alcohol	54 lb.
Zinc Oxide, Dry	29 lb.
Zine Dust	11 lb.
Pine Oil	5 lb.

Anti-fouling Paint

Gum Shellac	14 lb.
Denatured Alcohol	45 lb.
Zinc Oxide, Dry	14 lb.
Indian Red (Iron Ox	ide) 15 lb.
Mercuric Oxide	8 lb.
Pine Oil	9 lb.

Anti-fouling Paint, shown above, is used with the Anti-corrosive Paint shown above.

Anti-corrosive Paint

Zinc Oxide, Dry	19 lb.
Venetian Red, Dry	9 lb.
Silica	9 lb.
Rosin (WW Grade)	15 lb.
Solvent Naphtha	38 lb.
Manganese Linoleate	13 lb.
Coal Tar	5 lb.
The state of the s	

Anti-fouling Paint

Zinc Oxide, Dry	24 lb.
Asbestine, Dry	7 lb.
Silica	8 lb.
Cuprous Oxide	15 lb.
Mercuric Oxide	4 lb.
Rosin (WW Grade)	25 lb.
Solvent Naphtha	34 lb.
Pine Oil	4 lb.
Coal Tar	6 lb.

The steel decks should be primed with red lead paint and finished with two coats of one of the following deck paints:

Gum Shellac	
Wood or Denatured Alcohol	
Venetian Red (Iron Oxide)	
Chrome Green, Dry	
Dron Black Dry	

Bilge and Tank Paints Black Flexible Paint

Petroleum Residuum	34	lb.
Rosin	7	lb.
Petroleum Spirits	29	lb.
Coal Tar Naphtha	6	lb.

Black Deck Paint

Lampblack, Dry	4 lb.
Spar Varnish	44 lb.
Petroleum Spirits	5 lb.
Paint Drier	18 lb.

Grav Deck Paint

Zine Oxide, Dry	33	lb.
Lampblack, Dry	6	lb.
Ultramarine Blue, in Oil	1/4	lb.
Spar Varnish	74	lb.
Paint Drier	1	lb.

Red Deck Paint

Red Lead, Dry	10	lb.	
Indian Red (Iron Oxide), Dry	25	lb.	
Aluminum Stearate	2	lb.	
Lampblack, in Oil	2	lb.	
Spar Varnish	44	lb.	
Paint Drier	18	lb.	

Note: Since this paint contains red lead it can be applied directly on the steel deck; that is, no red lead primer is necessary.

Black Anchor Chain Paint

Gilsonite	7 lb.
Rosin	5 oz.
Petroleum Residuum	21 lb.
Solvent Naphtha	47 lb.

Green Anchor Chain Paint

Chrome Green, in Oil	10	Ib.
Red Lead, Dry	10	Ib.
Aluminum Powder	5	Ilr.
Asphaltum Varnish	4	gul.
Boiled Linseed Oil	• >	gal.
Spar Varnish	52	gal.
Petroleum Spirits	2	gal.
Paint Drier	1/2	gal.

Shellacs are used to brighten up wood work on marine vessels. The following are ten gallon formulas:

Orange	Red	Green
(clear)	Shellac	Shellae
24 lb.	27 lb.	27 lb.
55 lb.	48 lb.	53 lb,
-	17 lb.	************
Protection		15 lb.
*********	ter incom	15 11.

Black Acid-Resisting Paint

	~ *****	
Petroleum Residuum	20	lb.
Paving Asphalt	15	lb.
Lampblack, Dry	5	Ib.
Beeswax	21/2	lb.
Petroleum Spirits	39	lb.
Paint Drier	51/2	lb.

Bituminous Enamel	Cobalt Paint Prier	
Petroleum Residuum 80 lb. Paving Asphalt 10 lb. Asbestos Fiber 5 lb.	Cobalt Acetate 5 lb Rosin Ester Gum 15 lb Raw Linseed Oil 19 lb Petroleum Spirits 39 lb).
Paving Asphalt 10 lb.	Rosin Ester Gum 15 lb).
Asbestos Fiber 5 lb.	Raw Linseed Oil 19 lb).
Note: This product must be heated for	Petroleum Spirits 39 lb).
application.		
the Particular of the Particul	A 1 - 1/ 37 1-1	
Potable Water Tank Paint	Asphaltum Varnish	
Marie Water rank rank	Paving Asphalt 35 lb. Manganese Resinate 7 lb. Litharge 1 lb. Raw Linseed Oil 5 lb. 5 oz Petroleum Spirits 39 lb.	
Metallic Brown, Dry 40 10.	Manganese Resinate 7 lb.	
Indian Red, Dry 15 16.	Litharge 1 lb.	
Zinc Oxide, Dry 8 10.	Raw Linseed Oil 5 lb. 5 oz	5.
Silica 5 ib.	Petroleum Spirits 39 lb.	
Metallic Brown, Dry 40 lb. Indian Red, Dry 15 lb. Zinc Oxide, Dry 8 lb. Silica 8 lb. *Amberol Varnish 54 lb. Petroleum Spirits 3 lb. Paint Drier 3 lb.		
Petroleum Spirits 5 10.	Damar Varnish	
Paint Drier 5 ib.		
*Amberol Varnish for Above Formula.	Batavia Damar Gum 47 lh Turpentine 22 lh Petroleum Spirits 21 lh).
Raw Tung Oil 35 lb.	Turpentine 22 lb).
Petroleum Spirits 39 lb.	Petroleum Spirits 21 lb).
Amberol Gum No. 226 10 lb. Raw Tung Oil 35 lb. Petroleum Spirits 39 lb. Cobalt Drier 34 lb.	And the second s	
and the state of t	Copper Paint for Wood Bottoms	
Black Tank Paint	0 0 11	
Petroleum Residuum 12½ lb.	Denatured Alcohol 50 lb Zinc Oxide, Dry 16½ lb Indian Red, Dry 16½ lb Cuprous Oxide 8 lb Pine Oil 9 lb). `
Litharge 134 lb.	Zina Ovida Dry 1814 II). `
Red Lead 1¼ lb.	Indian Red Dry 161/2 It). `
Rosin (D Grade) ¼ lb.	Currous Oxido	٠.
Lampblack, Dry 5½ lb.	Pina Oil 0 11). `
Boiled Linseed Oil 12 lb.	I life Off	٠.
Spar Varnish 14 lb.	1 11 77 11 777 1	
Petroleum Residuum 12½ lb. Litharge 1¾ lb. Red Lead 1¼ lb. Rosin (D Grade) ¼ lb. Lampblack, Dry 5½ lb. Boiled Linseed Oil 12 lb. Spar Varnish 14 lb. Damar Varnish 4 lb. Petroleum Spirits 32½ lb.	Anti-Fouling Waterline Paint	
Petroleum Spirits 32½ lb.	Gum Shellac 13 lb. Denatured Alcohol 5 gal Pine Oil 3 gal Crude Rubber 1 oz. Gasoline 2 gill Zinc Oxide, Dry 5 lb. Lampblack, Dry 4 lb. Mercuric Oxide 4 lb. Turpentine 2 lb.	
Section of the sectio	Denatured Alcohol 5 gal	
Brown Tank Paint	Pine Oil 3 gal	
Drown rank rame	Crude Rubber 1 oz.	
Metallic Brown, Dry 40 lb.	Gasoline 2 gill	s
Litharge 2 lb.	Zinc Oxide, Dry 5 lb.	
Zine Oxide, Dry 16 16.	Lampblack, Dry 4 lb.	
Zine Chromate, Dry 2 15.	Mercuric Oxide 4 lb.	
Damar Varnish 40 10.	Turpentine 2 lb.	
Deint Deien 15 lb		
Metallic Brown, Dry 40 lb. Litharge 2 lb. Zinc Oxide, Dry 16 lb. Zinc Chromate, Dry 2 lb. Damar Varnish 46 lb. Interior Varnish 11 lb. Paint Drier 15 lb.	White Water Paint	
	Zinc Oxide, Dry 24 lt Whiting, Dry 48 lt Plaster Paris 24 lt	•
Primer for Bituminous Enamel	Whiting, Dry 48 lb	
Trinidad Asphalt 53 lb. Petroleum Spirits 63% gal.	Plaster Paris 24 lk). 1
Petroleum Spirits 6% gal.	Pulverized (Hide) Glue 4 lk	``
***	Plaster Paris 24 lk Pulverized (Hide) Glue 4 lk Ultramarine Blue, Dry 1 oz	z.
Bituminous Enamel	Note: Mix 8 lb. of the above mixt	
Paving Asphalt 52 lb. Trinidad Asphalt 15 lb. Rock Asphalt 15 lb. Rosin (Dark Grade) 1 lb. Portland Cement 17 lb. Slacked Lime 2½ lb.	in one gallon of water.	ure
Trinidad Asphalt 15 lb	in one ganon or water.	
Rock Asphalt 15 lb.	The same of the sa	
Rosin (Dark Grade) 1 lb.	White Enamel	
Portland Cement 17 lb.	Titanox B, Dry 72 lb Spar Varnish 28 lb Damar Varnish 29 lb Pine Oil 6 lb	١.
Slacked Lime 21/6 lb.	Spar Varnish 28 lb	
Note: This product must be heated be-	Damar Varnish 29 lb	
fore applying.	Pine Oil 6 lb	
	Ultramarine Blue, in Oil 1 oz	ž.
Point Di		
Paint Driers Manganese Resinate 10 lb. Damar Gum 10 lb. Litharge 2 lb. Raw Linseed Oil 8 lb. Petroleum Spirits 49 lb.		
Manganese Resinate 10 lb.	Gray Enamel	
Damar Gum 10 lb.	Titanox B, Dry 60 lb).
Litharge 2 lb.	Lampblack, in Oil 2 lb).
Kaw Linseed Oil 8 lb.	Titanox B, Dry 60 lb Lampblack, in Oil 2 lb Interior Varnish 53 lb Petroleum Spirits 9 lb).
Petroleum Spirits 49 lb.	Petroleum Spirits 9 lb).

Red Enamel		
Indian Red, Dry	4	0 lb.
Interior Varnish	5	55 lb.
Paint Drier		9 lb.
Petroleum Spirits		6 lb.
Outside White Pai	$_{ m nt}$	
Zinc Oxide, in Oil	50	lb.
Basic Sulphate White Lead	,	
in Oil	50	lb.
Blanc Fixe, in Oil	12	lb.
Asbestine, in Oil	6	lb.
Raw Linseed Oil	4	gal.
Petroleum Spirits	$\frac{3}{4}$	
Paint Drier	1/2	gal.
Ultramarine Blue, in Oil	1	oz.

Red Lead, Dry Silica 40 lb. Raw Linseed Oil 61/4 gal. Petroleum Spirits % gal. Paint Drier % gal.

Red Lead Paint

85

lb.

Light Gray Paint lb. Zinc Oxide, Dry Blanc Fixe, Dry 34 lb. Graphite, Dry 2 lb. Lampblack, in Oil 1 oz. Ultramarine Blue, in Oil 1 oz. Raw Linseed Oil 6% gal. Petroleum Spirits 1 gal. Paint Drier ¾ gal.

The formulas shown require that the pigments in oil be stiff pastes. The percentages of raw linseed oil present are within the limits shown:

	%	Lii	ise	ed C)
	•	in	Pa	iste	
White Lead (Carbonate)		8	to	10	
White Lead (Sulphate)		8	to	10	
Zinc Oxide		8	to	18	
Titantium Pigment B			15		
Chrome Green		33	to	35	
Chrome Oxide, Green		29	to	31	
Chrome Yellow		24	to	26	
Metallic Brown		22	to	24	
Lampblack		65	to	80	
Raw Sienna		45	to	55	
Burnt Sienna		40	to	50	
Raw Umber		35	to	45	
Burnt Umber		30	to	50	
Yellow Ochre		30	to	40	
Magnesium Silicate		20	to	30	
Venetian Red		20	to	25	

Black Marine Paint	
Carbon Black	15 lb.
Kaolin	25 lb.
Barytes	35 lb.
Boiled Linseed Oil	10 lb.

Rec	l Paint	
Indian Red Barytes Whiting Linseed Oil		5 lb. 1 lb. 1 lb. 2 lb.
Japan Drier Mixing Varnish		6 oz. 5 lb.
Su	rfacer	
Varnish		1 gal.
Brown Japan Silex (Fine)		1 gal. 8 lb.

Ship Bottom Paints 1. For Wood Bottoms

In any formulation, the object should be, first, to produce a mixture which will best serve the purpose and, second, to obtain the mixture at the lowest cost. The work requires a knowledge of a wide range of materials, their chemical and physical properties, and their cost. It also requires a knowledge of paint manufacturing operations, especially those to which the equipment on hand is adapted. Formulating is not an exact science any more than is the prescribing of medicine by the physician. One important difference between the physician writing a prescription and a paint technologist pre-scribing a paint formula is that the latter is also thinking about the cost.

The requirements of a paint for wood bottoms are comparatively simple and The corrosion problem easy to meet. does not enter, and consideration of a possible chemical or physical conflict with a priming paint does not enter. The object is to produce a paint, the film of which will brush (or spray) on easily, will dry quickly, will be resistant to water erosion and yet sufficiently softened by the water to permit the toxic elements to go into solution. There are several ways of approaching the problem which can best be illustrated by used formulas.

Formula No. 1

TOTHING ITO.		
Iron Oxide	18	Ib.
Silica	. 5	lb.
Copper Cyanide	13.5	lb.
Spar Varnish	7.25	gal.
Pine Tar Oil	.625	
Paint Drier	.23	gal.
"Tar Acid Oil"	.30	gal.
Mineral Spirits	.25	gal.

(Comment: The above formula will doubtless "dry" in about four hours because the spar varnish, which usually requires about twelve hours to dry, has been overloaded with the added driers. The dried film will be glossy and apparently hard, but it will probably not dry hard because of the excessive pine tar oil. The toxicant, copper cyanide, is regarded as only fairly toxic. This fact, together with the fact that a spar varnish film usually disintegrates under sea water and fouls readily, suggests that the film will not prevent barnacle fouling for a longer period than two or three months.)

Following is a formula which has given very good service:

NT	_	c
LY	() ₋	- 2

Blanc Fixe	40	lb.
Mercuric Oxide	5	lb.
Paris Green	7.5	lb.
Gum Shellac	20	lb.
Denatured Alcohol	5.9	gal.
Pine Oil	2.5	gal.

(Comment: The above formula is typical of shellac type paints. This paint will be effective about six months on a wooden bottom. It probably will not stand long storage satisfactorily, the nature of the pigment being such as to suggest a very hard sediment forming).

suggest a very hard sediment forming).
The U. S. Navy used a formula similar to the above.

No. 3

Zinc Oxide	165 lb.
Indian Red	165 lb.
Cuprous Oxide	75 lb.
Gum Shellac	162 lb.
Alcohol	500 lb.
Pine Oil	90 lb.

2. For steel bottoms.

In successfully formulating paints for steel bottoms the maximum ingenuity of the paint technologist is required. There are wide variations of opinions among men engaged in this work and each opinion is based, more or less, on experience in research. In designing paints for exposure to atmospheric elements there are certain fairly well established rules as to pigment-vehicle ratios by weight and by volume. For an oil paint for outdoor exposure, the pigment should be about 60 per cent by weight, and about 29.5 per cent by volume, of the paint. No such rules have been, or can be, established for ship bottom paints. Such ratios vary with each change in the vehicle, and there are an almost infinite number of such changes that can be made. The setting of high and low limits for the variants is apparently useless.

Before considering the varnish type of paints, which general type constitute the bulk of ship bottom paints used in Amer-

ica, the hot plastic paints, such as are used extensively in European countries, will be considered. Following are formulas used about twelve years ago by one of the European Navies.

Anti-corrosive Paint

Rosin	26.5	lb.
Benzol	26.5	lb.
Ozokerite	5	lb.
Iron Oxide	42	lb.

Anti-fouling Paint

moraouning rame		
Rosin	38.6	lb.
Stearin	14.7	lb.
Benzol	12.8	lb.
White Lead	7.4	
Verdigris	9.6	
Arsenic	13.2	lb.
Mercuric Oxide	3.7	lb.

To illustrate the varnish type ship bottom paints, two sets of paints used by the United States Navy are shown.

Anti-corrosive Paint

Formula No. 1

One Gallon Formula

Zinc Oxide		3.05	lb.
Zinc Dust		1.1	lb.
Gum Shellac		.425	lb.
Yacca Gum		.44	lb.
Alcohol		.8	gal.
Pine Oil		.067	
	No. 2		

Coal Tar		47.5	lb.
Rosin		145	lb.
Coal Tar Naphtha		380	lb.
Magnesium Linoleate		129	lb.
Venetian Red		93	lb.
Zinc Oxide		186	lb.
Silica		93	lb.
Beeswax	*	3.3	lb.

Anti-Fouling Paint Formula No. 1

One Gallon Formula

Zinc Oxide		1.65	lb.
Indian Red		1.65	lb.
Mercuric Oxide		.75	lb.
Gum Shellac		.815	lb.
Yacca Gum		.89	lb.
Alcohol		.76	
Pine Cil		.125	gal.
NI o	0		_

Coal Tar	132.6	lb.
Rosin	202.0	lb.
Coal Tar Naphtha	228.0	lb.
Pine Oil	74.0	lb.
Zinc Oxide	212.0	lb.
Silica	82.0	lb.

83.0 lb.

Asbestine

Cuprous Oxide Mercuric Oxide 112.0 lb. 45.0 lb.

Although commercially made phenolformaldehyde condensates have not proved satisfactory in undersea water exposure, there apparently is considerable merit to a varnish from such resin when the resin is made simultaneously with the varnish. These varnishes comprise the vehicle of the ship bottom paints and are made in reflux condensers. Typical of the process is the following:

Place 90 lb. of phenol, 108 lb. of 40% solution of formaldehyde, 90 lb. of water and 54 lb. of lead acetate in a reflux condenser and boil about 30 minutes. Add 720 lb. of rosin and continue heat until excessive foaming starts. Remove the reflux and continue heat until foaming ceases and at same time blow air through the mixture. Cool and add 108 gal. of coal tar naphtha.

The varnish is mixed with pigments to form anti-corrosive and anti-fouling

paints.

Anti-Corrosion and -Fouling Paint

Yacca Gum	1.6	lb.
Alcohol	1.32	gal.
Pine Oil	1.9	gills
Petroleum Spirits	1.9	gills
Zinc Oxide	1.2	lb.
Silica	1.2	lb.
Blanc Fixe	1.2	lb.
Zinc Dust	0.3	lb.
Paris Green	0.6	lb.
Mercuric Oxide	1.4	lb.

Paints for Ship Bottoms Formula No. 1

2.5 parts of wood tar, 2.0 parts of oxide of iron, 1.0 part of turpentine resin, 2.0 parts of lead acetate. Wood tar is preferable to coal tar, since the latter is not as resistant towards the corrosive action of sea water.

No. 2

1.0 parts of lead arsenate, 1.0 parts of Scheele's green (copper arsenite), 8.0 parts of ochre, 5.0 parts of turpentine resin, 3.0 parts of coal tar, 2.0 parts of Bakelite, 5.0 parts of oil of turpentine and 5.0 parts of white spirit.

No. 3

The so-called "Lucchini Paint": 30.0 parts of galipot (white resin produced from fir), 20.0 parts of turpentine resin, 2.5 parts of mercury arsenate, 20.0 parts of red arsenic, 30.0 parts of wood tar, 5.0

parts of manganese dioxide and 15.0 parts of oil of turpentine.

No. 4

600.0 parts of asphaltum or pitch, 480.0 parts boiled linseed oil, 120.0 parts of graphite, 120.0 parts of arsenic-copper oxide and 640.0 parts of coal tar oil.

No. 5

48.0 parts of coal tar, 383.0 parts of tar oil, 146.0 parts of turpentine resin, 130.0 parts of manganese linoleate, 3.5 parts of beeswax, 93.0 parts of Venetian red, 93.0 parts of infusorial earth and 187 parts of zine oxide.

No. 6

133 parts of coal tar, 288 parts of tar oil spirits, 20 parts of turpentine resin, 74 parts linseed oil, 21 parts of zinc oxide, 82 parts of infusorial earth, 83 parts of magnesium silicate, 112 parts oxide of copper and 145 parts mercury oxide.

Ship Bottom Paints

An anti-corrosive paint is prepared from 145 parts of oiticien fatty acids, 120 parts shellae, 390 parts alcohol, 80 parts pine oil, 5 parts drier, 170 parts of iron oxide, and 160 parts of zine oxide.

The anti-fouling composition given is, 145 parts of oiticica fatty acids, 129 parts shellac, 430 parts alcohol, 80 parts pine oil, 5 parts drier, 170 parts of iron oxide, 160 parts of zinc oxide, and 100 parts red oxide of copper, and 40 parts of yellow oxide of mercury.

To prepare these paints the shellac is dissolved in some of the alcohol, the pine oil added, and then the pigments ground in, using a ball mill. When perfectly smooth, the fatty acids, mixed with the pine oil and the remainder of the alcohol, are added. The same method is used in the case of the anti-fouling composition except that the yellow oxide of mercury is not ground in but only mixed.

After storage these products appear very thick, but spread easily under the brush to give a thick flexible film. A little diluent can be added if necessary.

The cobalt, lead and manganese salts of the fatty acids of oiticica oil, which are in the nature of driers, are prepared by converting the acids into soluble soaps and precipitating these by the acetate of the appropriate metal. The precipitate is carefully washed and dried in a carefully regulated oven in a current of earbon dioxide to prevent oxidation.

Paints for Glazing and Coloring Ceramics
Pigments (glass-powder, colored with
metal oxides)

Thinner: Linseed Oil Wood Oil

5 lb. 3 lb.

Artificial Mother-of-Pearl British Patent 426,554

Dibasic lead phosphate (PbHPC₄) prepared by adding phosphoric acid to a warm solution of a lead salt, if used in the form of very fine crystals, produces glistening or iridescent effects. The salt may be obtained in a very fine state of division by precipitation in presence of a water-soluble organic compound, and preferably under slightly acid conditions. Thus 24 liters of a solution of lead nitrate (3.3 kg. dissolved in 10 liters of water) are mixed with 4.8 liters of distilled water and 24 liters of 95 per cent alcohol; 2.6 liters of phosphoric acid (12 kg. of concentrated acid plus 50 liters of 90 per cent alcohol) are added all at once. The use of this lead phosphate in materials which are to be submitted to treatment with formaldehyde (e.g., casein products) is advantageous in that the salt is not affected by formaldehyde.

Enamel Opacifier British Patent 427,850

A fine white powder is obtained by heating at 1000° for several hours an intimate mixture of titanium dioxide 33.1, antimony pentoxide 44.5, and zine oxide 22.4%.

Pearl or Fish Scale Essence German Patent 603,487

Formula No. 1

a. Scales of Uklei Fish

b. {n-Propyl Acetate { 1.5 kg. } Nitrocellulose } 100 kg. 150 l. }

Treat a with b in a stirring-machine for 30 minutes, pour off the upper suspension, and repeat the same treatment of a two more times. Now the scales are free of fish-silver, the suspension containing about 1500-2500 g. of this substance.

No. 2

a. Herring Scales, Norwegian 100 kg.
b. { Ethyl Propionate 150 l. } Nitrocellulose 1.5 kg.

As in No. 1. Yields in fish-silver are quantitative, viz. 700-1000 g. crude material.

No. 3

a.	Astrachan Scales	100 kg.
b.	Ethyl Acetate	150 1.
	{ Ethyl Acetate { Nitrocellulose	3 kg.

Work as in No. 1, stirring two times for 20 minutes. 1200-1600 g. crude fishsilver can be obtained by centrifugal separation of the suspension.

No. 4

a.	Uklei	Scales	(from	Lake

Scutari, Albania) 1 kg. b. Ethyl Acetate 1.5 l.

c. { Acetyl Cellulose 25 g. { Alcohol, a little to dissolve completely.

As in No. 1 in smaller proportions. Should yield 1.4 liter suspension with 0.4% dry fish-silver.

Protective Coating for Hydrofluoric Acid Containers

Beeswax 1 oz. Paraffin Wax 4 oz.

Electric Lamp Coating U. S. Patent 1,941,990

The lamps are coated with a paste of Kaolin 50 g.
Guignet's Green 200 g.
Cadmium Sulphide 50 g.
Boric Acid 160 g.
Sodium Silicate (d. 1.015) 1000 cc.

Coating Lacquer for Fabrics

Coating Lacquer for Fabrics

Nitrocellulose Wet (5.6 sec.) 12 g.
Diamond "K" Linseed Oil 8 g.
Crude Crepe Rubber (Light) 8 g.
Ethyl Acetate 10 g.
Butyl Acetate 10 g.
Alcohol 5 g.
Toluol 47 g.

Heat crude rubber in linseed oil until dissolved. Cool and dilute with part of toluol. Add to remainder of formula after nitrocellulose is dissolved.

Rubberized Cloth Varnish Formula No. 1

Shellac 5 oz. Alcohol 95 oz. Gives high gloss.

No. 2

Shellac		1	kg.
Ammonia	(28%)		kg.
Water		30	kg.

Two coats of this must be applied to get good adhesion. The finish is semi-glossy. These varnishes are applied by a velvet covered brush or roller.

Waterproofing Brick Walls

Walls can be waterproofed by applying a coat of solution made by dissolving 1% lb. of paraffin in each gal. of mineral spirits used as a solvent. Use steam to melt rather than a free flame.

Moisture proofing Compositions Canadian Patent 352,183

Moistureproofing compositions consist of (parts by weight): Formula No. 1, paraffin 85, refined carnauba wax 10, rubber 5; (No. 2) paraffin 65, rubber 5, candelilla wax 30; (No. 3) paraffin 75, rubber 5, gum damar 20; (No. 4) paraffin 40, rubber 5, carnauba wax 40, ester gum 15; (No. 5) paraffin 60, rubber 5, carnauba wax 20, gum damar 15; and (No. 6) paraffin 55, rubber 4, candelilla wax 25, hydrogenated castor oil 16 parts.

Jute Waterproofing French Patent 763,402

Asphalt Bitumen Coal Tar Coal Tar Pitch Linseed Oil, Boiled Sand. Fine	5 2	lb. lb. lb.
Sand, Fine Bordeaux Resin	15	

Straw Lacquer Waterproofing Italian Patent 267,765

Cellulose Nitrate	10	oz.
Butyl Acetate	20	oz.
Benzol	48	oz.
Butyl Alcohol	7	oz.
Paraffin Wax	2	oz.
Camphor Oil	8	oz.
Butyl Ether	5	oz.
-		

Waterproofing Compound and Paint Vehicle

U. S. Patent 1,965,042

Three gallons of china-wood oil is raised to a temperature of about 240° C.; at this temperature 12 grams of manganese borate is added with rapid stirring. The temperature is maintained for a period not exceeding about fifteen minutes, but preferably from one to two minutes. In order to quickly cool the oil and also to partially dilute it, about 1 gallon of water white kerosene is added.

The temperature of the mass will thus be reduced to about 175° C. and when this temperature is attained 1½ pints of carbon tetrachloride is gradually added by introducing the same preferably near the bottom of the vessel. The rate of introduction of carbon tetrachloride is such that from 1–2 minutes are required for this step of the process. When the carbon tetrachloride has been introduced and the temperature has been reduced sufficiently, for example, to about 100° C, any desired quantity of diluent such as kerosene or solvent naphtha is added.

This forms a solution of waterproofing material which when applied to stone, brick, masonry and the like penetrates the pores of the same and coats the surface of the material to which it is applied, efficiently protecting it from the elements such as rain, sea water, salt water air, heat and frost. The coating is not substantially acted upon by alkalies or acids and forms a colorless waterproofing material which remains effective for many years.

Waterproofing Composition Belgian Patent 400,446

The composition contains carbon tetrachloride or carbon disulphide 200 cc., paraffin 150 g., rubber 8 g., and naphthalene 50 g. per liter.

Waterproofing Composition U. S. Patent Serial Number 513,225

A waterproofing composition which comprises forming a mixture of from 285 to 290 parts of water, 12 to 16 parts of sodium silicate and 9 to 10 parts of oleic acid and then stirring into this mixture approximately 300 parts of comminuted cumar resin (melting point about 230° to 245° F.) while maintaining the liquid at a temperature above 90° F. and not to exceed substantially 160° F.

Moisture and Greaseproof Coating

rolling 100. 1		
Gelatin	5.4	oz.
Sulfonated Oil	2.7	oz.
37% Formaldehyde Solution	1.4	oz.
Glycerin Monophthalate		
Ester	4.5	0.2

No. 2 In another specific formula, to each 100 oz. of a vehicle containing 10% alcohol add the following:

Gelatin	7.2	02.
Glycerin	3.6	oz.

37% Formaldehyde Solution 1.8 oz. Glycerin Monophthalate Ester 3.6 oz.

No. 3

In a third specific example add to each 100 oz. of vehicle:

Gelatin 5.8 oz. 37% Formaldehyde Solution 1.4 oz. Glycerin Monophthalate Ester 5.8 oz.

The two latter formulas, however, do not have the full effectiveness of the first in producing moisture-resistant and

greaseproof coatings.

In preparing the composition, when alcohol is employed in the vehicle, it is kept separate from the remaining constituents of the mixture until a late stage in the formation thereof. The gelatin is dissolved in a portion of the water, and, if desired, may be mildly acidulated, for example, with acetic acid. The flexibility-imparting agent, if any is used, is added to the aqueous solution of gelatin, suitably after admixture with or solution in a small amount of water, although this is not necessary. The formaldehyde solution is diluted with water. The diluted formaldehyde solution is then added, or, in its place, suitable proportions of a solution of hexamethylenetetramine, or alum or the like may be employed. The alcohol is diluted, suitably with an equal amount of water, and then The glycerin added to the mixture. phthalate ester or other ester employed is then dissolved in part or all of the remaining quantity of water, neutralized, for example, with ammonium hydroxide, and incorporated in the mixture.

Waterproof Finish

	Formula No. 1	No. 2
Tornesit	20 g.	20 g.
Methyl Abietate	12 g.	16 g.
Cumar V	12 g.	24 g.
Indian Red	25 g.	
Titanium Dioxid	le —	40 g.

Waterproofing Fibrous Materials U. S. Patent 1,965,630

One thousand pounds of pulp fiber dry weight is mixed in an ordinary paper mill beater with about 20,000 pounds of water. To this is added about 300 pounds of alkaline filler such as calcium carbonate, 15 pounds of ammonium resinate (dry weight) is then added in the form of an aqueous solution containing 500 pounds of water. 15 pounds of alum

are then added, which immediately reacts with the carbonate to form theoretically 3½ pounds of precipitated alumina. Instead of adding this alum to the beater, the alum solution may first be neutralized with ammonia or other alkali, and the precipitated alumina added to the beater with the size. The hydrated aluminum oxide will combine in the beater with the ammonium resinate to form a compound which coats the fibers in the beater and which will size the paper when the pulp is dried.

Another method of operation is as fol-

lows:

The carbonate filler, or other filling material, is mixed with water in a tank to a concentration of about 20% solids to which mixture is added an aqueous solution containing ammonium resinate to the extent of about 1 pound of the dry resinate to 100 pounds of filler. To this may be added 1 pound of alum to each 100 pounds of filler along with sufficient ammonia or other alkali to neutralize it

and precipitate the alumina.

This separately treated filling material containing sizing ingredients may be added to the paper stock in the beater, in the Jordan chest, in the machine chest, or at the wet end of the paper machine. This treatment produces a paper containing individually sized filler particles, that is, each particle thereof is coated individually with size. The paper stock in the beater may be sized by the use of ammonium resinate and alumina. If this is done, the result is a paper with fibers and filler particles individually sized with the same sizing materials. Or the paper stock may be first sized with any sodium resinate and sufficient alum to acidify the fibers, whereupon and later, the ammonia sized filler material is added thereto in the beater, machine chest, Jordan, and so forth, whereby a paper is produced having its fibers individually sized by the use of sodium resinate while its filler particles are individually sized with ammonium resinate and alumina. Since the ammonium resinate is somewhat more expensive than sodium resinate, this latter procedure offers some saving in cost over treating both fibers and filler with ammonium resinate.

In general, in the final mixture of paper fibers and filling material, there must be no alkalinity derived from soda. There will be none in the mixture resulting from the practice of this invention because any alkalinity produced by the ammonium resinate disappears on drying of the paper. This produces a neutral

and sized paper.

3 lb.

With the present processes using sodium resinate, it is not possible to fully size a heavily loaded paper containing from 20% to 30% filler even if the filler is not alkaline. By the use of this process, however, any kind of filling material can be sized. In order that ammonium resinate may properly function as a sizing material there should always be present enough excess ammonia or other alkali, to form sufficient alumina when reacting with alum to form a resinate of alumina, but it is immaterial how this ammonium hydrate is produced.

Waterproofing Composition U. S. Patent 2,022,405

Refined Paraffin Wax	4 lb.
Paracoumarone Resin	2 lb.
White Beeswax	1 lb.
Aluminum Palmitate	4 lb.

The above ingredients being blended together and dissolved in a composite solvent of xylol and carbon tetrachloride in the proportions of about three parts by volume of xylol to one part by volume of carbon tetrachloride, and the amount of solvent being such that about 23/4 ounces of the above composition is contained in each gallon of solution.

Fireproofing Materials French Patent 774,089

An antiseptic fireproofing composition for wood, paper, etc., contains, e.g., ammonium orthophosphate 5 grams, sodium tetraborate 2.5 grams, and ammonium chloride 2.5 grams.

Exterior Primer

Pigment	67	lb.
Vehicle	33	lb.
Pigment:		
Titanox B	37.1	lb.
White Lead (Carbonate)	37.1	lb.
Asbestine	24.8	lb.
Litharge	1.0	lb.
Vehicle:		
Archer-Daniels No. 635	64	lb.
Mineral Spirits	26	lb.
*VM-1367	8	lb.
2% Liquid Cobalt Drier	2	lb.

*VM-1367: Heat 15 gal. china wood oil with 75 lb. low acid ester gum to 565° F. Remove from fire and let rise to 585° F., hold for 5 minutes and check with 25 lb. ester gum. Thin at 400° F. with 15 gal. mineral substite. spirits.

Painting Primer German Patent 608,738

Zinc Oxide Ochre Linseed Stand Oil Linseed Oil Varnish The above is thinned with:	30 g. 30 g. 14 g. 21 g.
Linseed Oil Varnish	3 g.
Benzine	9 g.

Exterior Wood Primer		
Pigment	66	lb.
Vehicle	34	lb.
Pigment:		
Titanium-Barium Pigment	34	lb.
White Lead (Carbonate)	26	lb.
Metronite	40	lb.
Vehicle:		
Bodied Linseed Oil	13	lb.
Blown Linseed Oil	5	lb.
Raw Linseed Oil	27	lb.
20-gal Ester Gumwood		
Oil Varnish	20	lb.
Mineral Spirits	32	lb.

Priming Paint from Hardened Paint German Patent 607,554

Dissolve old paint in following:

Drier

Butyl Alcohol	50	lb.
Xylol	10	lb.
Benzol	10	lb.
Toluol	10	lb.
Ethyl Acetate	. 5	lb.
Ether	. 5	lb.

Galvanized Roof Primer

Dry Red Lead	10	lb.
Boiled Linseed Oil	91/2	gal.
Turpentine	17/8	gal.
Drier	1/8	gal.

Galvanized Roof Finish

	Dry Red Lead	5	lb.
٠	*Carbon Black Paste	31	lb.
	Boiled Linseed Oil	61/4	gal.
	Turpentine		gal.
	Drier		gal.

*The carbon black paste referred to in this formula is 18.0% carbon black and 82% boiled oil.

Paste Paint L, White Lead

Basic Carbonate White	е	
Lead	28.4	lb.
Raw Linseed Oil	3.88	lh.

30 THE CHEWICA	II POINTULAILI
Paste Paint TLZ, Titanox-Lead-Zinc Titanox B 9.8 lb.	White Exterior Bakelite Enamel (Yacht White)
Basic Carbonate White Lead 7.6 lb.	Pigment 40 lb. Vehicle 60 lb.
Zinc Oxide, Lead-Free 4.33 lb. Raw Linseed Oil 3.88 lb.	Pigment:
Spot Priming Paint	Basic Carbonate White Lead 40 lb. Titanium-Barium Pigment 40 lb. Titanium Oxide 20 lb.
Paste Paint TLZ (above) 1 gal. Raw Linseed Oil 1 gal. Turpentine 0.28 gal.	Vehicle:
Turpentine 0.28 gal. Drier 0.05 gal.	*Varnish XV-4430 60 lb. †Varnish XV-5922 20 lb. Mineral Spirits 20 lb.
Under Coat Paint	Drier:
Paste Paint L or TLZ (above) 1 gal. Raw Linseed Oil 0.51 gal. Turpentine 0.62 gal. Drive 0.04 gal.	Lead 2.5 g. per gallon enamel, as naphthenate
Turpentine 0.62 gal. Drier 0.04 gal.	Cobalt 0.15 g. Manganese 0.05 g.
Finish Coat Paint	*Varnish XV-4430: Bakelite Resin XR-2963 100 lb. China Wood Oil 20 gal.
Paste Paint L or TLZ (above) 1 gal. Raw Linseed Oil 1 gal. Turpentine 0.12 gal. Paint Drier 0.06 gal.	Bakelite Resin Ar-2963 100 15. China Wood Oil 20 gal. Body Q Linseed Oil 30 gal. Lead Acetate 2 lb. Mineral Spirits 34 gal. Dipentene 5.5 gal. Procedure:
Paint Drier 0.06 gal.	Place the Bakelite, the China wood oil and
Tropical Roofing Paint Paste White Lead 100 lb. Non-setting Red Lead 10 lb. Lamp Black in Oil ½ lb. Raw Linseed Oil 3 gal. Boiled Linseed Oil 1 gal.	Place the Bakelite, the China wood oil and 10 gallons of the linseed oil in the kettle. Heat to 560° F. in one hour. Add the remaining 20 gallons of linseed oil. The temperature will drop to about 450° F. Reheat to 520° F. Add the lead acetate. Cool quickly with the aid of water spray to 450° F., and thin with the mineral spirits. †Varnish XV-5922:
Turpentine or White Spirit ½ gal. Terabine Driers 1 pt.	Bakelite Resin XR-2963 100 lb. China Wood Oil 7.5 gal. Body Q Linseed Oil 2.5 gal. Lead Acetate 2.5 lb. Lead Carbonate 1.25 lb. Mineral Spirits 15 gal.
A proportion of hard drying outside quality varnish may be added if desired. Thin out this paint to the desired consistency with equal parts of raw linseed oil and turpentine. Where the paint must be cheapened, barytes, china clay, slate powder, or ochre is incorporated as an extender.	Procedure: In 50 minutes heat the Bakelite and China wood oil to 450° F. In an additional 18 minutes raise the temperature to 540° F. Add the linseed oil and the driers. Let the temperature drop to 450° in about 20 minutes, and thin with the mineral spirits.
Priming Structural Paint	Lead Titanate Exterior Paints
Formula No. 1	Formula No. 1
Dry Basic Lead Chromate 15½ lb. Raw Linseed Oil 5 pt. Turpentine 2 gills Liquid Drier 2 gills No. 2	Lead Titanate 1000 lb. Raw Linseed Oil 252 lb. China Wood Stand Oil 28 lb. Lead-Manganese-Cobalt Drier 8 lb. Mineral Spirits 42 lb.

 $\begin{array}{cccc} \text{Dry Basic Lead Chromate} & 15\frac{1}{2} & \text{lb.} \\ \text{Boiled Linseed Oil} & 5 & \text{pt.} \\ \text{Turpentine} & 1 & \text{pt.} \end{array}$

These paints weigh about 21 pounds per gallon and the non-volatile portion contains about 30% by volume of pig-

ment,

. Formula No.	1		
Lead Titanate	10	000	lb.
Raw Linseed Oil	9	252	lb.
China Wood Stand Oil			lb.
Lead-Manganese-Cobalt	Drier	8	lb.
Mineral Spirits			lb.
No. 2			

No. Z		
Lead Titanate	400	lb.
Basic Carbonate White		
Lead	400	lb.
Asbestine	100	lb.
Silica	100	lb.
Pour Tinggood Oil	200	11-

China Wood Stand Oil Cobalt Naphthenate Mineral Spirits	52 10.8 65.9	lb.
No. 3		
Lead Titanate Basic Carbonate White	400	lb.
Lead	400	lb.
Zinc Oxide	200	lb.
Raw Linseed Oil	$\frac{200}{324}$	lb.
Kettle Bodied Linseed Oil	524	10.
	01.0	11.
(Viscosity Z)	21.6	10.
Lead-Manganese-Cobalt	00.4	12
Drier	20.4	
Mineral Spirits	40.7	16.
No. 4		
Lead Titanate	400	1b.
Titanox-B	400	lb.
Zinc Oxide	200	lb.
Raw Linseed Oil	400	lb.
Kettle Bodied Linseed Oil		
(Viscosity Z)	26.4	lb.
Lead-Manganese-Cobalt		
Drier	25.1	lb.
Mineral Spirits	50.1	
No. 5		
Lead Titanate	200	lb.
Titanox-B	200	lb.
Basic Carbonate White		
Lead	200	lb.
Zinc Oxide	200	lb.
Asbestine	100	lb.
Silica	100	lb.
Raw Linseed Oil	466	lb.
Kettle Bodied Linseed Oil	400	10.
(Viscosity Z)	29.6	115
Lead-Manganese-Cobalt	25.0	TI).
Drier	29.2	1h
	58.4	
Mineral Spirits		
*This type is of special interest base for house paint tints.	st for	use as

Fire Retarding Interior Whitewash

a base for house paint tints.

1. Mix about 120 lb. of spent carbide residue with water to a creamy consistency.

 Mix 2½ lb. of rye flour thoroughly with ½ gal. of cold water, and then thin with 2 gal. of boiling water.

3. Dissolve 2½ lb. of common salt in 2½ gal. of hot water.
Mix (2) and (3), then add (1), and stir until well mixed.

Exterior Weatherproof Whitewash Formula No. 1

- 1. Mix about 120 lb. of spent carbide residue with water to a creamy consistency.
- Dissolve 2 lb. of common salt and 1 lb. of zinc sulphate in 2 gal. of boiling water.

3. Provide 2 gal. of skimmed milk. Pour (2) into (1), then add (3), and stir well.

No. 2

 Mix about 15 lb. of spent carbide residue to a creamy consistency with water.

 Dissolve 1 lb. of carbonate of soda in ¼ gal. of boiling water.

3. Soak in cold water for at least 8 hr. ¼ lb. of common glue and 1 lb. of rice flour; and then thoroughly dissolve the glue mixture in ¾ gal. more water in a double boiler. Mix (1) with (2), then add (3).

No. 3

- 1. Mix about 12 lb. of carbide residue to a creamy consistency with water.
- 2. Dissolve 4 oz. of white rosin in 12 fluid oz. of boiled linseed oil.
- Beat 6 lb. of whiting in 1 gal. of skimmed milk.
 Mix (2) with (1) while hot, add (3).

Hints for Special Uses

Alum added to whitewash prevents its rubbing off. Flour paste will also prevent rubbing off, but when this is used, zinc sulphate must be added as a preservative.

Molasses causes lime to penetrate wood and plaster better. One pint of molasses to 5 gallons of whitewash is generally considered sufficient. A solution of silicate of soda or water glass, one part to ten parts of whitewash, makes what is commonly referred to as a "fire-proof cement" of whitewash.

By adding 1 pound of cheap bar soap dissolved in 1 gallon of boiling water, to every 5 gallons of whitewash, a more or less gloss finish can be obtained.

A fire retardant whitewash, of a type used extensively by the U.S. Lighthouse Board, is made according to this formula:

- Mix about 60 lb. of spent carbide residue with water to a creamy consistency.
- 2. Dissolve 1 peck of salt in warm water.
- 3. Add (2) to (1) and mix.
- 4. Boil 3 lb. of ground rice in water to to a thin paste.
- 5. Dissolve 1 lb. clear glue in hot water.
- 6. Provide ½ lb. of powdered Spanish whiting.
- 7. Mix (4), (5), and (6) together and add to mixture (3). Mix well and let stand for several days.

Keep the wash thus prepared in a kettle or portable furnace, and when used put it on as hot as possible with a painter's brush or whitewash brush.

Cold Glaze for Wall Tiles

Υ		T
Laco	uer	Base

acquer Base	
a. Shellac	8 oz.
Turpentine, Thick	5 oz.
Alcohol	35 oz.
b. Sandarac	14 oz.
Turpentine, Thick	6 oz.
Alcohol	35 oz.
Mix 10 oz. of a with	

Mix 10 oz. of a with 12 oz. of b

To this lacquer base add finely powdered pigments, as to color

Lamp Black	(Black)
Ultramarine or Paris Blue	} (Blue)
Chrome Yellow Zinc Yellow	(Yellow)
or Ochre Chrome Green	(Green)
Chrome Red	(Red)
or Cinnabar	(1000)
Lithopone	(White)

(Grind Pigment with a small part of the lacquer solution; thin later with the rest to needed consistency.)

Floor Finish

(Permanent, Scratch-free) Clear (Natural) Finish:

Formula No. 1

Castor Oil	1 qt.
Boiled Linseed Oil	½ gal.
Paraffin Wax	3¼ lb.
High-Flash Naphtha	3 qt.
Gasoline	1½ gal.
Varnolene	1 gal.

Mix the oils and wax and heat until the wax is molten. Add the varnolene, naphtha and gasoline slowly in the order mentioned.

No. 2

Dark Finish

Castor Oil	1	qt.
*Gilsonite Cook	1	gal.
Paraffin Wax	3	Ĭb.
High-Flash Naphtha	1	qt.
Gasoline	11/2	gal.
Varnolene	1	gal.

Heat oil and wax until molten, add the gilsonite cook and proceed as above.

*Gilsonite Cook:

Chadhie Cook.		
Gilsonite	5	lb.
Kellogg Varnish Oil	11/2	gal.
High Flash Naphtha	11/4	gal.
Heat gilsonite and oil to 270° C.	(52	0° F.)
Let cool and thin with naphtha.		

Any shade may be obtained by intermixing clear and dark finish. Apply by flowing on the freshly scraped floors, distribute and rub in lightly with rags. Permit to dry for at least 48 hours. This finish actually impregnates the floor and will not wear off. It has a velvet sheen and a slight slip, is easy to keep clean and is very resistant to moisture.

Varnish for Naval Aircraft

Materials:

Bakelite BR-254	50	lb.
Bakelite XR-4036	50	lb.
Castor Oil (Refined)	4.33	lb.
China Wood Oil	33	gal.
Mineral Spirits	27	gal.
Xylol	4	gal.
Dipentene	4	gal.

Lead-Cobalt-Manganese Naphthenate Driers

Procedure:

Heat the oil and the Bakelite resins together to 310° F. in 25 minutes, and hold at that temperature for half an hour. Heat to 450° F. in 20 minutes and hold for 20 minutes. Remove from the fire, add the thinners, the castor oil and sufficient drier to give 12 grams cobalt, 15 grams manganese and 160 grams lead as metal.

Airplane Varnish

The naval aircraft factory has developed a formula for satisfactory bituminous varnish which is used for airplane hulls or other parts exposed to salt water or salt spray. This formula is as follows:

Aluminum	Powder	2	lb.
Bituminous	Primer	1	gal.

Coating for Aluminum or Brass Nitrocellulose 5 g. Amyl Acetate 55 cc. Alcohol 40 cc.

Aluminum Powder Paste U. S. Patent 2,002,891

Aluminum, Flaked Stearic Acid, Powdered		oz.
Aluminum Stearate		oz.
Naphtha	40	oz.
Grind together until homoge	eneor	s.

Preparing Aluminum for Enamel

The best method of cleaning aluminum castings, so the finish will adhere tenaciously, is to use the sandblast. Smooth

aluminum surfaces are of such character that an ordinary first coat of finishing material will not adhere to them satisfactorily, even when they are clean. The sandblast will leave the surface slightly etched and will aid the first coat in sticking to the metal permanently.

If sandblasting is impractical, about all that can be done is to thoroughly wash the castings with naphtha or some other solvent for grease, and dry them

thoroughly with clean cloths.

In other instances it may be satisfactory to bake the castings for a short time at 400 or 500° F., just before finishing them, to burn off any oil or grease. It is not advisable to use caustic cleaning solutions with aluminum, because the metal is so easily attacked and dissolved by this chemical.

Another method is as follows: Immerse them in a 20% solution of acetic acid until all oil and grease is removed or neutralized. Then rinse in a vat of clear hot water and allow castings to drain and dry. Do not wipe them. Spray or brush as soon as the moisture has disappeared.

Bronzing Liquid

Celluloid Scrap	3 oz.
Amyl Acetate	12 oz.
Benzine	28 oz.
Denatured Alcohol	24 oz.

This solution is mixed with sufficient dry gold bronze to make a smooth working paint and the resulting paint must be used at once as it is apt to turn greenish and thicken to a jelly on standing.

Bronze Painting Tinctures

	. (Wate	er.		90	oz.
А.	a. \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \	ho l		10	oz.
	b. Isingl	ass or	Mirror		
	0 1 7			2 .	. 7

Gelatin as desired
Add to this colloidal solution with stirring:

c. Bronze Powder sufficient to suit. B. for a and b take:

Potash-Water	Glass	10	oz.
Gum Arabic		10	oz.
Water		40	oz.

C. or Thick Gum Arabic Solution with a little ox-gall.

Paints for Copper

Copper, bronze, or brass gutters and flashings, as well as copper or bronze screening, are apt to cause bad yellowishgreen stains on light- or white-painted

houses, owing to the washing off of corrosion products. Exposure tests indicate that one of the best ways to paint copper or bronze surfaces is to wash off any grease, using gasoline or turpentine. The surface should be roughened slightly with sandpaper, and a priming coat composed of 11/2 to 2 pounds of aluminum powder to 1 gallon of aluminum mixing varnish applied, followed by the desired color coat. Weathered copper or bronze screening should be thoroughly dusted, and then given two coats of a thin black paint. Some of the best grades of black auto top dressings, which are free from asphalt, but are essentially thin, water-resistant, carbon black enamels, make excellent screen enamel.

Cable Lacquer British Patent 397,554

Cellulose Acetate	12	oz.
Triacetin	12	oz.
Mineral Oil		
(b.p. 330-390° C.)	0.8	oz.
Acetone	50.2	oz.
Toluol	10	OZ.
Alcohol	10	OZ.
Diacetone Alcohol	5	oz.

Electrolytic Condenser Coating British Patent 419,927

Acetone	137.8	ee,	
Amyl Acetate	125.0		
Phenol-Formaldehyde Resin	39.9	g.	
Graphite (99%)	42.5	g.	

This is baked on aluminum for 24 hours at 100° C, and 2 hours at 170° C,

Electrical Wire Lacquer British Patent 410,576

Cellulose Acetate	100	oz.
Tetrachlorethane	100	02.
Alcohol	20	oz.
Triacetin	3	OZ.

Adhesiveness may be increased by incorporating tale and opacity by zine oxide.

Wash for Galvanized Iron before Painting

a.	Denatured Alcohol	60	fl.	oz.
	Toluol	30	fl.	OZ.
	Carbon Tetrachloride	5	fl.	02.
	Commercial Concentrate	ed		
	Hydrochloric Acid	5	fl.	oz.

b. Copper Acetate 6 oz. Water 1 gal. c. Copper Nitrate Crystals 2 oz. Copper Chloride Crystals 2 oz. Ammonium Chloride Crystals 2 oz.

Commercial Concentrated
Hydrochloric Acid 1/6 pt.
Water 1 gal.

Solution a will cut grease as well as etch. If the metal is not free from grease, solutions b and c must be preceded by a grease-removing operation.

Treatment of Galvanized Sheets for Painting

A simple and inexpensive way to treat new galvanized sheets before painting is to use ordinary vinegar, either sponged or brushed on. Vinegar rather thoroughly removes the slick film usually found on newly galvanized sheets. It does not, however, etch the surface like some other treatments. After the vinegar has been applied and allowed to remain on the sheets for five minutes or so, it should be wiped and then the surface of the sheet allowed to completely dry before paint is applied.

Another somewhat similar treatment is the use of two or three per cent acetic acid solution at a temperature of about 130° F. If it is possible to dip the sheets, or articles made from the sheets, in this solution, allow them to remain there for about ten or fifteen minutes. After removal, they should be thoroughly rinsed and allowed to thoroughly dry.

Still another, even more practical, although perhaps a little more costly, method of obtaining a clean and etched surface is to apply, with an oil-free brush, and allow to remain for about ten minutes, an acidified solution made up as follows:

Denatured Alcohol 50 fl. oz. Toluol 35 fl. oz. Hydrochloric Acid 5 fl. oz.

This solution should be prepared only as required for immediate use. After the reaction is complete and the surface is thoroughly dried, wash or rinse with clean water to remove any soluble salts that may have formed. Then, allow the sheets to thoroughly dry again before applying paint. This treatment is especially effective if the procedure outlined above is carefully followed.

It should be particularly noted that with each of the three methods outlined, it is important that the galvanized surface should be thoroughly dry before painting. A film of moisture between the paint and sheet would cause very poor adherence.

Painting Galvanized Iron

Excellent paint adherence on galvanized surfaces may be obtained by cleaning with the following solution:

Alcohol	65	lb.
Toluol	35	lb.
Muriatic Acid (Commercial		
Concentrated)	5	lb.
Carbon Tetrachloride	10	lb.

This treatment should be followed by a cold rinse after the material has dried.

Lacquer for Hot Water	Contain	ers
Lacquer Linseed Oil	250	g.
Milori Blue	15	g.
Gilsonite	120	g.
Albertol Resin (116° mp	.) 40	g.
Thick Linseed Oil	40	g.
Manganese Hydroxide	2.5	g.
Cobalt Drier	1.25	g.
Toluol	500	g.

Iron "Lacquer"

Gilsonite Asphalt	20	kg.
Manila Copal		kg.
Lampblack	3	kg.
Toluol	50	

Iron Protective Paint

Formula No. 1

Lampblack (Ground in Oil)	90.7	oz.
*Asphalt Varnish	68.1	
Linseed Oil, Raw	68.1	oz.
Japan Drier	$^{2.0}$	oz.

No. 2

740. 7	
Lampblack	27 oz.
Silica	58 oz.
Red Lead	10 oz.
Graphite	5 oz.
*Asphalt Varnish	Sufficient
Grind together until smo	ooth.
*Turpentine Asphalt in Linseed Oil	1 part 1 part

Primers for Light Metal Alloys

Owing to high coefficient of expansion and contraction with temperature changes, a primer is needed that will be sufficiently flexible not to be ruptured by expansion and contraction. A zinc chromate paint is recommended for this purpose, a specimen formula being:

Zinc Chromate	40	lb.
Neutral Red Oxide of Lead	80	lb.
Boiled Linseed Oil	60	lb.
Pure Turpentine	16	lb.
Strong Japan Driers	4	lb.

Another priming paint found to be satisfactory is made from:

Dry Lampblack 65 lb. Linseed Oil 15 lb. 10 lb. Pure Turpentine

Driers according to type and quality. The primer should be allowed 50 to 60 hours to dry and harden before applying subsequent coatings.

Polished Metal Lacquer

Nitrocellulose Wet (15-20 sec.) Rezyl No. 468-2 (50% So-	10	g.
lution) Dibutyl Phthalate Butyl Acetate Butyl Alcohol Butyl 'Cellosolve'' Toluol Xylol	10 2 10 8 10 35 15) tá tá tá tá tá

Preparing Magnesium Alloys for Painting

To prepare the surface of magnesium alloys so that paint will adhere, it is recommended that the alloy be first immersed in the following:

Sodium Dichromate 1.5 lb. Concentrated Nitric Acid 1.5 pt. gal.

In a new solution, only 15 seconds are needed. This time increases to two

minutes for an old solution.

After rinsing and drying, the proper primer should be used, containing inert pigments or, for example, zinc chromate. For interior work, a minimum of two coats (total) paint should be used; for exterior work, a minimum of four coats.

Care and Preservation of Bronze Statues

Statues, tablets, medals, especially those standing in the open, require careful treatment and protection from the conditions tending to their corrosion. Of cleansing reagents, water only is permissible with, perhaps, a small quantity of soap extract. Bronze which has become black by long exposure may be restored to its original gold color by washing with water to which a little ammonia is added, using a brush with bristles, no wire brush.

As protective coating, a mixture of beeswax and turpentine is considered the best, it affords considerable protection to bronze from atmospheric attack and gives a pleasing appearance, besides drying rapidly. Applied three times a year it will safeguard a statue to a high degree from corrosion and deterioration even in an exposed position. A mixture of lanolin and paraffin is not quite as good as it does not dry as rapidly and is therefore liable to collect dust.

Heatproof Rust Protective Coatings

Kerosene and pitch cannot be used as binders as they become too soft even at 150-200° C. Natural asphalts, although brittle, give protection up to 250° C., acetyl cellulose up to 100° C. Only lean, not fatty binding agents rich in resins, should be used for such paints. As at 400° C. almost all binding agents are entirely disintegrated, the residues of the agents must be such that they leave a continuous, well adhering coat on the metal to be protected. Durophen, aluminum bronze, zinc dust with binders of this type give good results. Heatproof paints should never be sprayed on, as they have the tendency to spall off later, but brushed on, except zinc dust which may also be sprayed.

Rust Proofing

A good protective coat for metal articles during storage and transit is made by brushing on a solution of lanolin in white spirit or solvent naphtha. Equal weights solvent and landlin seem satisfactory and there is not much to choose between the two solvents. rather harder film is wanted, up to 5% ceresin wax can be added in the case of naphtha solutions; in the case of white spirit up to 10 per cent paraffin wax or up to 3 per cent ceresin wax. It is recommended that the white spirit be of the B.E.S.A. standard, i.e., B.P. 160° to 210° approximately and as to the lanolin, the results of practical tests show little difference between widely different grades.

7.8 lb. lanolin in 1 gal. white spirit give 1.9 gal. solution.

8.3 lb. lanolin in 1 gal. solvent naphtha give 1.9 gal. solution.

Crystal Coating on Steel

Sodium Nitrate 3 lb. Manganese Dioxide 3 lb. 8% Sulphuric Acid Solution 100 gal.

Protective Coating for Structural Steel

Coal-Tar Pitch	62.5	lb.
Benzol	25	1b.
Aluminum Bronze	12.5	lb.

90 lb.

Priming Structural Paint (Red Lead) Formula No. 1

a difference and a			
Dry Red Lead	20 lb.		
Raw Linseed Oil	5 pt.		
Turpentine	2 gills		
Liquid Drier	2 gills		
No. 2			
Red Lead Paste in Oil	20 lb.		
Raw Linseed Oil	3 pt.		
Turpentine	2 gills		
Liquid Drier	2 gills		

Finish for Steel Surfaces

Tornesit	20 g.
Linseed Oil, Crude, Boiled	10 g.
Indian Red	20 g.
Xylene	30 cc.
High-Flash Naphtha	40 cc.

First Coat Structural Steel Protective

Blue Lead, Paste in Oil	100	lb.
Raw Linseed Oil	23/8	gal.
Turpentine or Mineral		•
Spirits	$1\frac{3}{4}$	gal.
Drier	1/4	gal.

Top Coal Structural Steel Paint Pigment:

C.P. Chrome Orange

Magnesium Silicate	10	lb.
Vehicle:		
Raw Linseed Oil	80	lb.
Spar Varnish	10	lb.
Liquid Paint Drier	10	lb.
Paint:		
Above Pigment	70	lb.
Above Vehicle	30	lb.

Red Lead for Bridges

Red Lead	40	lb.
Iron Oxide (95%)	40	lb.
Stand Oil	90	lb.
Raw Linseed Oil	12	lb.
Turpentine	20-40	lb.
Cobalt-Manganese Drier	1	lb.

Tin Can Coating U. S. Patent 2,009,776

A coating dough for producing a coating material comprises a mixture of 100 parts by weight of rubber solution containing approximately 30 parts by weight of rubber, approximately 15 parts by weight of adhesive ester gum, approximately 3 parts by weight of liquid petrolatum, and approximately 100 parts by weight of zinc oxide.

Tin Lithographing Varnish

Typical construction of this class of product is represented by the following formulae: 54 gal. pale amberol varnish, 34 gal. gum solution, 22 gal. pale mixing varnish, 8 lb. of white vaseline warmed and reduced with 2 gal. of mineral spirits.

The first component of the above blend, is—135 lb. amberol F7 light, 15 lb. WWX Rosin, 34 gal. pale China wood oil, 1½ gal. "Superior" linseed oil, 6 gal. bodied linseed (1½ hrs. at 600° F.), 10 lb. fused lead resinate, 1 ounce cobalt acetate, 8 gal. gum turpentine, 65 gal. mineral spirits.

The second component is a solution of ester gum in mineral spirits, using 12½ lb. of gum to each gallon of solvent.

The third component is 50 lb. ester gum, 3 lb. fused lead resinate, 10 lb. WWX Rosin, 50 gal. pale China wood oil, 50 gal. mineral spirits.

"Tornesit" Paints

First, a base solution is prepared, consisting of 33½ per cent Tornesit and 66½ per cent high-flash naphtha. To effect solution is a matter of a very few minutes, if the "Tornesit" is added to the solvent.

Second, a concentrated gum solution is made when required.

Third, the pigments are ground in the plasticizer, or if it is insufficient, some of the "Tornesit" base solution is used.

Fourth, if a brushing paint is required, the base solution is thinned to a "Tornesit" content of 21 per cent to 22 per cent by the addition of a solvent mixture consisting of two parts high-flash naphtha and one part xylol. If a spraying composition is desired, the base solution is thinned with toluol to a "Tornesit" content of 11 per cent to 12 per cent. It is advisable to ship even spray paints with a brushing viscosity and send the thinner separately. This helps to keep the pigments in good suspension.

Finally, the gum solution and pigment paste are added to the reduced solution and the mixture is stirred.

"Tornesit" paints may be applied by spraying, dipping, flowing, or brushing. A good film can be obtained by any of these methods.

Following is a brief outline of procedure to be followed, to obtain most satisfactory results in spraying and brushing:

Spraying

"Tornesit" solutions can be sprayed, producing a hard, durable, evenly distrib-

70 g.

uted film. With present equipment, the spraying viscosity is 40 centipoises, which is somewhat lower than the 75 centipoise spraying viscosity of lacquers.

If the "Tornesit" concentration is kept below 12%, no difficulty will be encountered from "spider-webbing." By the addition of softening agents, guns, and pigments, the solids content will be increased 30-40 per cent, depending, of course, on choice of ingredients.

Brushing

Brushing paints with as high as 57 per cent solids have been applied successfully. For this purpose, a working viscosity of about 250 centipoises is recommended. In brushing ''Tornesit'' paint, the surface should be well covered with a full brush, avoiding going over the painted area any more than necessary because of the rapid drying of the product. When bodied tung oil is used as the plasticizer in the priming coat, a second coat may be applied to an interior surface after six to eight hours. On exterior work, three to four hours is an ample drying period with the same priming coat.

"Tornesit" Paints

A formula used successfully on tank cars, structural steel and similar surfaces not subject to immersion contains Tornesit plasticized with a drying oil. China wood oil must be properly boiled to avoid wrinkling when a second coat is applied, but no wrinkling occurs with linseed oil. When properly formulated, "Tornesit" paint has good adhesion to metal. Examples of primers having good adhesion are:

	Form	nula	No. 1	No	. 2
"Tornesit"		20	oz.	20	oz.
Heavy-Bodied	Raw				
Linseed Oil		10	oz.	10	oz.
Cumar PlO				5	oz.
Iron Oxide		20	oz.	20	oz.
Silica		30	OZ.		
Xylol		70	0 Z .	70	oz.

A finish coat used successfully on steel contained:

"Tornesit"	20	oz.
Heavy-Bodied Raw		
Linseed Oil	10	OZ.
Indian Red	20	oz.
Xylol	30	oz.
High-Flash Naphtha	40	oz.

Formulæ containing improperly-bodied oils do not have good alkali resistance, but to withstand immersion in aqueous media, particularly those containing alkalies, formulæ such as the following have been quite successful:

Formula No. 1 No. 2

"Tornesit"	20 oz.	20 oz.
Methyl Abietate	12 oz.	16 oz.
Cumar V	12 oz.	24 oz.
Indian Red	25 oz.	
Titanium Dioxide		40 oz.

Finishes made to the foregoing formula containing iron oxide have withstood immersion in 5 per cent caustic soda for two months and in 5 per cent hydrochloric acid for three weeks, the use of iron in the pigment probably reducing resistance to hydrochloric acid.

Pliolite Varnish (Paper Coating) Pliolite Resin 15 g. Ester Gum Solution (4 # cut) 10 g. Trieresyl Phosphate 5 g.

Toluol

Paper Enamel U. S. Patent 2,000,453

Glue	20	oz.
Ammonium Hydroxide	2	oz.
Alcohol	4	oz.
Chromic Acid	11/2	OZ.
Water to make	1	gal.

Moisture Proof Paper Lacquer British Patent 412,687

Ozokerite	1-2 oz
Dibutyl Phthalate	25-50 oz.
Nitrocellulose	50-75 oz.
Lacquer Solvent	to suit

Paper Watermarking Fluid U. S. Patent 2,021,141

C. 10. W. 1000-110		
Canada Balsam	8-20	lb.
Turpentine	5-17	lb.
Colorless Mineral Filler	8-25	lb.
Castor Oil	12-30	lb.
Borax Solution (1%)	sufficient	to
emulsify above liquid	s.	

Water to thin to working consistency.

Rubber Paints

British Patents 407,038 and 417,912

Preparation of Solution "B"

Raw crepe rubber is masticated on a rubber mill, using warm rollers, until the rubber runs coherently round the rollers. Keeping the rubber still milling, 2½ per cent of cobalt linoleate (6 per cent metallic cobalt content) is then added. When the cobalt linoleate is completely dispersed in the rubber, the mixture is

taken off the mill and immediately transferred to a solution mixer, and churned up with an equal weight of white spirit, until a homogeneous mass is formed. This is then poured into drums and is ready for use. The solution should not be kept at a lower concentration than 50 per cent, as there is a tendency for thinner solutions to reduce still further in viscosity and to lose some of their properties.

Preparation of Paint

To prepare a paint, the rubber solution is mixed to the oil with sufficient white spirit to make a medium, which when mixed with the necessary pigments, will form a suitable paste for grinding. Any of the usual pigments and fillers can be incorporated. The ground paste is then thinned with further white spirit to brushing consistency.

As examples of up-to-date formulæ for rubber paints, the following are suggested:

Flat Paints

Formula No. 1

Lithopone	150	lb.
Yellow Ochre	1.5	lb.
Middle Chrome Yellow	1.5	lb.
Solution "B" (above)	20	lb.
Boiled Oil	10	lb.
White Spirit	30	lb.

No. 2

Lithopone	65	lb.
Titanium White	65	lb.
Asbestine	15	lb.
Solution "B"	20	lb.
Stand Oil	10	lb.
Liquid Driers (Lead .033;		
Cobalt .004)	1	lb.
White Spirit	30	lb.

No. 3

Ultramarine Blue	75	lb
Asbestine	25	lb
Solution "B"	28	lb
Boiled Oil	14	lb
White Spirit	60	lb

No. 4

Lithopone	100	lb.
Solution "B"	20	lb.
Ester Gum	10	lb.
White Spirit	50	lb.

No. 5

Lithopone	150	lb.
Stand Oil/Wood Oil (3/1)	10	lb.
Solution "B"	20	lb.
Liquid Driers	1	lb.
White Spirit	30	lb.

Ready-Mixed Gloss Paints

No. 6

Zinc Oxide	100	lb.
Pale Boiled Oil	62.5	lb.
Solution "B"	25	lb.
Terebene	2	lb.
White Spirit	10	lb.
No. 7		
Zinc Oxide	50	lb.
Titanium White	50	lb.
Pale Boiled Oil	62.5	lb.
Solution "B"	25	lb.
Terebene	2	lb.
White Spirit	10	lb.
No. 8		
Lithopone	80	lb.
Zinc Öxide	20	lb.
Pale Boiled Oil	30	lb.
Solution "B"	15	lb.
Terebene	.1	lb.

No. 9

20 lb.

White Spirit

White Lead	100	1b
Pale Boiled Oil	30	1b
Solution "B"	12	lb
Terebene	1	lb
White Spirit	6	$^{\mathrm{1b}}$
White Spirit	6	11

Cheap Rubber Paint

Molten Rubber	100	oz.
White Spirit	100	oz.
Terebene	12	oz.
Cobalt Terebene	12	oz.
Red Ochre	100	oz.

The defects of molten rubber as a paint vehicle may be obviated by using it in conjunction with oil. That is to say, the varnish is made up partly of molten rubber and partly of linseed oil. A paint made up on a varnish of this description prepared by "cooking up" the oil and rubber together (in the proportion of 50/50) in the presence of driers and thinning with solvents—appears to have good ageing properties and to yield a film which does not readily crack.

¥	Molten Rubber Varnish	140	0.00
		7.40	OZ.
	Terebene	5	oz.
	Red Ochre	100	oz.
	* Molten Rubber * Linseed Oil + Driers White Spirit		oz.
	* {Linseed Oil + Driers	35	oz.
	White Spirit	7.0	0.7

Rubber Water Paint

Glue Solution		25	oz.
Casein Solution) ×	25	oz.
Latex		30	oz.
Lithopone		100	oz.

Drying oils can, if desired, be incorporated with the above, and for some

purposes are an advantage, but tend to discolor the paint more rapidly.

Distempers can also be satisfactorily prepared by using a rubber solution (as used for the oil paints). The solution readily emulsifies with a glue solution, with which the pigments can be incorporated.

The following is an example of this

type of distemper:

type or disterriper.	
Glue Solution	20 oz.
* Rubber Solution	16 oz.
Water	25 oz.
Lithopone	100 oz.
{Milled Crepo * {Cobalt Linoleate White Spirit	8 oz.
* Cobalt Linoleate	0.2 oz.
White Spirit	8 oz.

Rubber Frosting Varnish

The addition of rubber solution to china wood oil gives a frosting varnish which will give the desired effect in a more regular manner than when china wood oil is used alone. The rubber solution containing cobalt linoleate is suitable for this purpose.

20 oz.
10 oz.
1 oz.
10 oz.
10 oz.
0.25 oz.
10 oz.

Rubber Flat Paint

Aubber Solution		91	oz.
Milled Crepe		1	
Rubber		0Z.	
Lead Linoleate	1	oz.	
White Spirit	40	oz.	
Stand Oil		10	OZ.
Cobalt Linoleate		0.25	oz.
Lithopone		150	oz.
White Spirit		40	oz.

Rubber Gloss Paint

* Rubber Solution	25	oz.
Pale Boiled Oil	$62\frac{1}{2}$	oz.
Terebene	2	oz.
Zinc Oxide	100	OZ.
White Spirit	10	OZ.
Modified rubber solution co	ntaini	ng co
balt linoleate (as pre	eviously	√ de

scribed).
Solution as above, after blowing with air.

* $\begin{cases} \text{Milled Crepe (including} \\ 2\frac{1}{2}\% \text{ Cobalt Linoleate,} \\ \text{White Spirit } 12\frac{1}{2} \end{cases}$

* Rubber Resin Varnish	25 oz.
Stand Oil	50 oz.
Terebene	5 oz.

Cobalt Linoleate	1 oz.
Zinc Oxide	100 oz.
White Spirit	40 oz.
* {Rubber Resin {White Spirit	162% oz. 81% oz.

Rubber Lacquer

THE THE THE THE THE	
Nitrocellulose, Wet	
(15-20 Sec.)	5.0 g.
Staflex Oil (Plasticizer)	2.5 g.
Ethyl Acetate	10.0 g.
Butyl Acetate	10.0 g.
C.D. Alcohol	10.0 g.
Toluol	62.5 g.

Rubber Repairing Lacquer (For Galoshes)

Alcohol 240	cc.
a. Nigrosin (Alcohol-Soluble) 2	g.
Nigrosin-Base BT 50	g.
b. Benzol (90%) 180	cc.
$b. \begin{cases} \text{Nigrosin-Base BT} & 50 \\ \text{Benzol } (90\%) & 180 \\ \text{Acetone, Technical} & 200 \end{cases}$	cc.
Ma 250 as of this depote of solution	n ad

To 350 cc. of this dyestuff solution add Xylene, Technical 350 cc. Vinapas B.P. 50T 300 g.

Mix thoroughly, filter through a gauze filter.

Black Rubber Tire Paint

Ditton	ALCONOCA	A. A. C.	A CALALO	
Rosin				kg.
Turpentine				kg.
Shellac				kg.
Sandarac				kg.
Alcohol				kg.
Turpentine,	Venice			kg.
Carbon Bla	ck		to	suit

Elastic Covering French Patent 762,342

Viscose	15 - 30	
Rubber Latex	50 - 80	g.
Casein	70	
Water	45	
Sodium Silicate (36° Bé.)	25	
Hardwood Flour	70	
Asbestos Fibers	35	
Ochre, Uncalcined	40 - 60	g.

Rubber-Asphalt Lacquer

Crepe Rubber (Shredded) 5-10 oz. Benzol 90-95 oz.

Allow to soak over night and stir the next day until uniform,

Dissolve

Gilsonite 30-40 oz.

Benzol 60-70 oz. Run the rubber solution into the other

solution slowly while stirring.

Linoleum Preservative

Formula No. 1

Linseed Oil (Free from Mucous Substances)

No. 2

Caoutchouc, Crude, Soft	45	g.
Resin, Coumarone	15	g.
Spindle Oil, Refined	940	ġ.
Melt up together on water	bath.	

Linoleum Finish U. S. Patent 1,998,927

Glyceryl Phthalate	12.5	lb.
Toluol	48.1	lb.
Triethanolamine	0.9	lb.

Apply to uncured plastic linoleum body and keep at about 75° C. for 14 days.

Eggshell Enamel

Pigment	50	Ib.
Vehicle	50	lb.
Pigment:		
French Process Zinc Oxide	80	lb.
Celite No. ON-165	10	lb.
Titanium Dioxide	10	lb.
Vehicle:		
Kettle-Bodied Linseed Oil	60	lb.
Mineral Spirits	35	lb.
Liquid Cobalt Drier	5	lb.

Enameling over Varnish

First wash wood work; sandpaper; mix flat paint or enamel undercoat with a little enamel and brush it out thinly. While wet rub with pumice stone and then smooth coating with a brush. Only a small section may be done at a time. If coating sets too quickly add a little linseed oil.

Aluminum Lacquer

Beckosol No. 1, Solid	100	g.
Solvent Naphtha	100	g.
Chlorinated Rubber	20	g.
Xylene	70	ğ.
Cobalt-Siccative (1% Coba	lt) 5	ğ.
This lacquer is resistant to	benzol	

Analytical Weight "Lacquer" Bleached Shellac Alcohol, Pure 4 fl. oz.

Put in corked bottle; shake and allow to stand for a few days. Filter through fine filter paper.

Brushing Lacquer U. S. Patent 1,533,616

+ · · · · · · · · · · · · · · · · · · ·		
Alcohol	10	oz.
Ethylene Glycol	10	oz.
Amyl Acetate	5	oz.
Butyl Acetate	10	oz.
Ethyl Acetate	15	oz.
Benzol	15	oz.
Toluol	10	oz.
Xylol	10	oz.
Gasoline	10	oz.
Amyl Alcohol	5	oz.
Butanol	5	oz.
Crystal Lacquer		

Nitrocellulose, Wet (1/2 sec.)	8	g.
Tunguran "A" (Plasticizer)	9	
Furfural	12	g.
Butyl Acetate		g.
Ethyl Acetate	30	
Toluol	33	g.

Lacquer Thinner

and the second		
Toluene	50	cc.
Ethyl Acetate	18	cc.
Alcohol	12	cc.
Amyl Acetate	20	cc_{\bullet}

Cellulose Solution No. 1

Nitrocellulose (Dry Weight)		
(½ sec.)	25	g.
Alcohol	10.7	g.
Butyl Acetate	16.1	g.
Toluene	32.1	
Ethyl Acetate	16.1	g.

No. 2

Nitrocellulose (Dr	
(½ sec.)	35.8 g.
Butyl Acetate	24.8 g.
Toluol	24.2 g.
Ethyl Acetate	15.2 g.

Crystallizing Lacquer Thinner

Ethyl Acetate	1.5	g.
"Cellosolve"	0.5	g.
"Cellosolve" Acetate	0.5	g.
Methanol	0.5	g.
Toluene	. 7	g.

If using phthalic anhydride, make up solution in cyclohexanone, if using naphthalene dissolve in toluene. The resulting solution is stirred into the lac-quer. Variations are made by using mixtures of both, naphthalene and phthalic anhydride.

Crystallizing Lacquer Formula No 7

rormula No. 1		
Cellulose Solution No. 1		
(see above)	15	g.
Cellulose Solution No. 2		Ĭ
(gga aharra)	0.5	~

0.02222100123		
Cyclohexanone Ester Gum in Toluol	6.5	g.
(1:1, Weight)	2	œ
Tricresyl Phosphate	0.5	g.
A contacts	5	g.
Amyl Acetate	•	g.
Phthalic Anhydride, or Naph	- ,	
thalene Flakes	4	g.
No. 2		
Nitrocellulose (1/2 sec.)	4	g.
Nitrocellulose (100 sec.)	1.5	g.
Butyl Acetate	9.5	g.
Ethyl Acetate	9.5	g.
Cyclohexanone	8	g.
Butyl Propionate	9.5	g.
Toluene	2	
Methanol	$\frac{5}{3.25}$	g.
	9	g.
Thinner (see below)	-	g.
Ester Gum in Toluol (I:1)	7.0	g.
Phthalic Anhydride or	a =	
Naphthalene Flakes	8.5	g.
The phthalic anhydride is t	o be	dis-
solved in the cyclohexanon	e (heat
gently), then stir solution into l		
8//		
Lacquer for Electric Bu	ıllıs	
Nitrocellulose	20	œ
		g.
Butyl Acetate	0.5	g.
Acetone	50	g.

Spirit (Furniture)	Lacquer	
Shellac, Bleached	25	g.
Sandarac	8	g.
Turpentine	4	cc.
Alcohol, Denatured	100	cc.

30 5-10

Alcohol

Lithopone, optional

Alcohol, Denatured

or other Pigments.

Floor Paint Lacquer Formula No. 1

Rosin	100	g.
a. \ Wood Oil, Crude	60	g.
$a. \begin{cases} ext{Rosin} \\ ext{Wood Oil, Crude} \\ ext{Linseed Oil} \end{cases}$	40	g.
b. Zinc White	4	g.
c. Litharge Manganese Oxide-Hydrate	3	g.
Manganese Oxide-Hydrate	0.5	g.
a. Lacquer-Denzonne		
(White Spirit)	160	g.

Heat up a together to 180-200° C., then add b together with lime (to harden the oils). Heat up to 290° C., take off the fire. When temperature falls to 250-260° C., add c.

When cooled, thin with d.

-	-	-
- N	•	•

a. {Kopol No. 600 a. {Wood Oil, Crude b. Linseed Oil—"Standoil"	100 g	ŗ.
Wood Oil, Crude	70 g	ζ.
b. Linseed Oil-"Standoil"		•
Thick	30 2	ς.

160 g. c. Lacquer Benzoline d. Cobalt-Siccative, Liquid (1% Metal Content) 6-8 g.

Heat a to 280-290° C., then "quench" with b. When cooled to 180° C., add c, then d.

Floor Lacquer

Copal Ester	100	g.
Linseed Oil—"Standoil"	70	g.
Lead-Manganese Resinate	4	g.
Cobalt, Siceative		g.
Thinner	150	g.

Linoleum or Floor Lacquer

manufacture of a root arms que		
Nitrocellulose, Wet (1/2 sec.)	14	g.
Dewaxed Damar Gum So-		
lution (4# Cut)	12	g.
Paraplex 5-B Solution (80%		
by Weight) (Plasticizer)	12	g.
Dibutyl Phthalate	2	g.
Toluol	15	g.
Mineral Spirits	20	g.
Butyl Alcohol	10	g.
Butyl Acetate	5	g.
Butyl "Cellosolve"	10	g.

Hat Lacquer

Use 1.25 gal. of the damar lacquer shown below to 3.75 gal. of the second thinner although other thinners can be used.

A lacquer may be made from damar gum and nitrocellulose as follows: 12.5 gal. benzene; 12.5 gal. toluol; 50 lb. 5-sec. nitrocellulose; 10 gal. ethyl acetate; 8.75 gal. butyl acetate; 21.25 gal. dewaxed damar solution.

The yield is 67 gallons of lacquer. Put the five-second nitrocellulose in a 100 gal, barrel or drum and wet it down with the toluol and a low boiling petroleum lacquer thinner. After mixing them, add the ethyl acetate, butyl acetate, and de-waxed damar solution. Stir by hand with a wooden stick, or a power stirrer. The dewaxed damar solution is made quickly by grinding to about 10 mesh: 80 pounds of No. 1 Batavia or Singapore damar gum and adding it to-2.7 gallons of ethyl acetate and-6.43 gallons of petroleum benzene or cleaners' naphtha. Stir this mixture until it is in solution and then as the stirring continues add: 17 gallons of 200 proof alcohol, for cutting shellac. After adding the alcohol, a white waxy precipitate will be formed which will take from one to three days for settling out, depending upon the kind of alcohol used.

The lacquer just described is usually thinned with two parts of a suitable thinner to one part of lacquer before dipping hats into it. The hats are put on racks to dry before shaping on the hot block. A very agreeable non-poisonous thinner is made by mixing: 53% cleaners' naphtha; 15% butyl acetate; 24% No. I Special or other similar solvent; 6% butanol; 2% butyl lactate.

Marble Effect Lacquering German Patent 597,114

Marble effects are gotten by applying the following oil coating over a ground coating of lacquer and then spraying on immediately a very thin lacquer.

Paraffin Oil	40	0 oz.
Toluol	20	0 oz.
Alcohol	20	oz.
Ethyl Acetate	20	oz.
Pine Oil	20	0 oz.
Castor Oil		5 oz.

Non-Inflammable Lacquer

Cellulose Acetate Plasticizer Ethylene Dichloride Ethyl Acetate Alcohol Methyl "Cellosolye"	20 g. 20 g. 120 g. 30 cc. 20 cc.
Methyl "Cellosolve" "Cellosolve" Acetate	20 cc. 5 cc.

Pavement Lacquers Formula No. 1

roimula 100. 1	
Rosin, Pale	14 g.
Manila Copal	30 g.
Linseed Oil	22 cc.
Cobalt Linoleate Drier	1 g. 33 cc.
Benzoline	33 cc.
No. 2	
Alcohol	40 cc.
a. Alcohol Manila Copal	40 g.
b. "Galipot" in Alcoholic Solution (1.5:1)	
	20 cc.
(Rosin in Alcoholic So-	
c. Rosin in Alcoholic Solution (2:1)	20 cc.
Mix solutions a , b , c .	

Lacquer Plasticizer

Coconut Fatty Acids	2610	lb.
Sulphuric Acid (66° Bé.)	about 500	
Denatured Alcohol Caustic Soda (14° B		gal.
Caustic Soda (30° B		

Manipulation:

1. The coconut fatty acids must be saponified by boiling with excess of

strong caustic soda solution (30° Bé. or stronger) and with addition of considerable water after saponification to prevent solidification of the soap.

2. This soap solution is then decomposed with sulphuric acid, the resulting coconut fatty acids (now being free from neutral oil) are washed with hot water.

3. The fatty acids are heating in a lead lined pressure vessel at 20 to 25 pounds pressure with denatured alcohol and sulphuric acid to esterify to the ethyl ester of the mixed fatty acids. This operation is carried on until the free fatty acid test shows only 6-7 per cent, beyond which point it is uneconomical.

4. The remaining free fatty acids are then neutralized with a 14° Bé. caustic soda solution in a steel tank, allowed to settle over night and the mixed esters pumped off from the scapstock to the still for distillation.

5. The esters are distilled under 25-26 second vacuum at a temperature of 250-425° F. in a steel mill equipped with oil heat or with means for circulating the esters through a direct heater. The condensing equipment is equipped with a sight glass so that the first runs, which are dark in color, may be separated for addition to the next lot of acids to be esterified. When the distillate becomes pale yellow it is suitable for the finished product receiver. The finished product is bleached water white with Fuller's Earth and decolorizing carbon.

Lacquer Thinners Formula No. 1

T OTH	ши	TA O.	1		
Ethyl Acetate				15	oz.
Butyl Propiona	te			25	oz.
Toluol				60	oz.
	No.	2			
Ethyl Acetate				5	oz.
Butyl Propiona	te			10	OZ.
Fusel Oil				20	OZ.
Toluol				55	OZ.
Xylol				10	oz.
	No.	3			
Amyl Acetate				20	oz.
Butyl Alcohol				- 10	oz.
Methyl Alcohol				10	OZ.
Toluol				60	oz.
	No.	4			
Benzine				40	OZ.
Amyl Acetate				10	OZ.
Butyl Acetate				30	oz.
Acetone				20	oz.

Lacquer for Synthetic Plastics

The consistency of a particular lacquer is governed in the first place by the pro-

25 oz.

25 oz.

posed mode of application. In general,
spray lacquers contain 12 to 14 per cent
nitrocellulose; dipping lacquers contain
8 to 12 per cent nitrocellulose, and brush
lacquers contain 14 to 17 per cent nitro-
cellulose.

Solvent mixtures will also vary with the mode of application. A typical solvent mixture for cellulose lacquers comprises:

Lacquer Solvent

Ethyl Acetate	50	oz.
Butyl Acetate	20	oz.
Butyl Alcohol	5	oz.
Benzol	25	oz.

The following lacquer compositions are recommended for highly polished surfaces:

Formula No. 1

Butyl Acetate	40	oz.
Ethyl Acetate	10	0 Z .
Alcohol	25	OZ.
Benzol	10	oz.

Remainder nitrocellulose, including 10 per cent plasticizer (calculated on the nitrocellulose) such as dibutyl or diamyl phthalate.

No. 2	
	oz.
Butyl Alcohol 5	oz.
	oz.
Glycol Monoacetate 5	oz.

Lacquer Sealers Formula No. 1

	1.04	
Nitrocellulose (1/4 sec.) Wet	2.22	lb.
Thinner	1	gal.
Lacquer Thinner.		

Toluol 61 oz. Butyl Acetate

101001	OI OZ.
Butyl Acetate	26 oz.
Butyl Alcohol	13 oz.
No. 2	
Nitrocellulose (1/2 sec.)	.47 lb.
Nitrocellulose (40 sec.)	.93 lb.
Ester Gum	.93 lb.
Calcium Stearate	.93 lb.
Thinner	1 gal.
Lacquer Thinner:	

Coal Tar Naphtha Butyl Acetate

	Sealing Lacquer		
	Formula No. 1		
Celluloid	Scrap	10	oz.
Lacquer	Solvent	30	OZ.

60 oz.

40 oz.

Denatured Alcohol	10	oz.
Barium Sulphate	25	OZ.
Zinc Oxide	25	oz.
No.		
Cellulose Acetate	15	oz.
Methyl Acetate	5	oz.
Lacquer Solvent	30	oz.
Barium Sulphate	25	oz.
Chrome Yellow		oz,
No.	3	
Nitrocellulose	10	oz.
Ether	15	oz.
Alcohol		oz.
Barium Sulphate		oz.
Ochre		oz.
No.		0.24
	-	0%
Pyroxylin		
Lacquer Solvent	35	02.

Greater adhesion can be secured in above formulæ by addition of 3 % ester gum.

Barium Sulphate

Chrome Orange

Capsule or Tube Sealing Lacquers Formula No. 1

Celluloid Scrap Lacquer Solvent Alcohol, Denatured Lampblack	15 oz. 40 oz. 25 oz. 20 oz.
No. 2	20 02.
Cellulose Acetate	20 oz.
Methyl Acetate	5 oz,
Lacquer Solvent	50 oz.
Zinc Oxide	25 oz.
No. 3	
Nitrocellulose	15 oz.
Ether	22 oz.
Alcohol	38 oz.
Ultramarine Blue	25 oz.

Transparent Tube Lacquer Formula No. 1

Callada Samon

Centinoia Scrap	20	OZ.
Lacquer Solvent	50	oz.
Alcohol, Denatured	20	oz.
Butanol	8	OZ.
Soluble Lacquer Color	2	oz.
No. 2		

Nitrocellulose	18	oz.
Butyl Acetate	15	oz.
Lacquer Solvent	68	OZ.
Soluble Lacquer Color	2	oz.

Lacquer for Tennis Rackets

Manila	Copal	33	g.
Alcohol	(93-95%)	66	
	Oil Fatty Acid	1	cc.

Flexible Gloss Wood Lacquer	"Aquarell" Colors
Nitrocellulose, Wet (1/4 sec.) 14 g.	Pigments
Ester Gum Solution (4# cut) 20 g.	White:
Blown Castor Oil 4 g.	Whiting Finest, or China Clay.
Dibutyl Phthalate 3 g.	Pale Yellow:
Ethyl Acetate 10 g.	Pale Yellow Lake, or Yellow
Butyl Acetate 10 g.	Lake, Blended.
Butyl Alcohol 7 g. Toluol 32 g.	Yellow:
101doi g.	Yellow Lake, Martius Yellow,
Ethyl Cellulose Wood Lacquer	Ochre.
Ethyl Cellulose (Low Vis-	Pale Orange:
cosity) 8 g.	Orange Lake, Blended to get
Dewaxed Damar Gum Solu-	Lighter Colors.
tion (4# cut) 12 g.	Orange:
Dibutyl Phthalate 2 g.	Orange Lake.
Alcohol 10 g. Toluol 58 g.	Rosa (Pink):
"Cellosolve" Acetate 10 g.	Alizarin Lake, or "Echt-Rot"
	(Genuine-Red), Blended to Ob-
West West Income	tain Lighter Colors.
Flat Wood Lacquer	Red:
Nitrocellulose, Wet (¼ sec.) 12 g.	Alizarin Red, Martius Red.
Dewaxed Damar Gum Solu- tion (4 lb. cut) 10 g.	Pale Brown:
Ester Gum Solution	Terra di Siena, Blended
(4 lb. cut) 10 g.	Brown:
Blown Castor Oil 2 g.	Caput Mortuum (Iron Oxide).
Dibutyl Phthalate 1 g.	Dark Brown:
Halowax No. 1014 5 g.	Umbra, or Cassel Brown.
Ethyl Acetate 5 g.	Violet: Brilliant Violet Lake.
Butyl Acetate 15 g. Butyl Alcohol 7 g.	
Toluol 25 g.	Pale Blue: Blue Violet Lake, Blended.
Xylol 8 g.	Blue:
Maria de la companya del companya de la companya de la companya del companya de la companya de l	Blue Lake.
Flexible Barrel (Inside) Coating	Dark Blue:
a. Gilsonite Asphalt 50 g. Benzol 50 cc.	Dark Blue Lake.
	Pale Green:
b. Caoutchouc, Crude 5 g. Benzol 50 cc.	Green Lake, Blended.
	Green:
Prepare a in an iron-kettle with stirrer, if necessary, heat.	Green Lake.
Prepare b soaking cold for several	Gray:
days. Mix the two viscous solutions,	Black Lake, Blended.
pouring b into a, stirring vigorously.	Black:
Apply repeatedly, allowing each layer	Black Lakes.
to dry well.	The blending, to get paler shades, is
T-13- C-41- C- W. 1	done by mixing the lake or pigment with
Inside Coating for Wood Barrels	white chalk.
a. Yellow Wax 40 g.	The state of the s
Colophony 200 g.	Manufacture of "Aquarell" Colors
b. Iron Oxide 40 g.	(Water soluble, applied with brush)
c. Gypsum (Molding) 10 g.	Solution for binding of the pigments
Melt up a , then stir in b , finally c .	in the color-paste:
Apply liquid, hot mixture with a brush.	Formula No. 1
Towns	Gum Arabic 26 g.
Lacquer for Barrels	Water, Distilled 51.9 g.
Rosin 22 lb.	Glycerin (28° Bé.) 8 g.
Turpentine, Thick 4 lb.	Glucose Solution (1:1) 10 g.
Turpentine 4 lb. Alcohol 12 lb.	Beef-Gall, Prepared 4 g.
12 10.	Moldex or Other Preservative 0.1 g.

Dissolve gum powder in cold water, stir, then heat to get complete solution. Add preservative, then glycerin, glucose solution, beef-gall. Filter, when cooled, through a percolator-cloth. (See No. 2)

No. 2

Dextrin, White	40	g.
Water, Soft or Distilled	41.8	g.
Borax, Crystallized	2	g_{\bullet}
Glycerin (28° Bé.)	G	$\mathbf{g}.$
Glucose Solution (1:1)	10	g.
Moldex or Other Preservat	ive 0.2	g.
Moldex or Other Preservat		

Make dextrin paste in cold water, then warm to get clear solution, add preservative and borax, then glucose-solution and glycerin.

Add the amount of water lost by evap-

oration (also in No. 1).

Alkali and Acid Resisting Paints Formula No. 1

18 lb.
43 lb.
9 lb.
4 lb.
9 lb.
17 lb.
1 lb.

No. 2

No. 2	
Chlorinated Rubber	18 lb.
Toluol	45 lb.
Gutta-Percha Resin	11 lb.
Wood Oil Stand Oil	2 lb.
Amyl Acetate	6 lb.
Tetralin	5 lb.
Paint Graphite	11 lb.
Carbon Black	1 lb.

Fireproof Paints (for Wood)

Barium Sulphate	25	oz.
Zine White	1	oz.
Water	20	OZ.
Waterglass	25	oz.

Heat Sensitive Paints

Certain chemicals in form of paints can be employed for the detection, or determination, of temperature fluctuations of a surface. Thus, the double iodide of silver and mercury, which is yellow at ordinary atmospheric temperatures, is colored dark orange on heating, being brick red at a temperature of 70 to 80° C. The double iodide of copper and mercury is bright red at ordinary temperatures, turning chocolate brown at 70° C. and black at 100° C. If the heating of the paint films is not ex-

tended too far, the original color of the paint returns on being cooled back to ordinary atmospheric temperatures. A process recently patented in France employes a mixture of two substances, which react upon each other at elevated temperatures only, lead sulphide and barium superoxide. In a suitable carrier this mixture is black at ordinary temperatures, turning gray on heating. This change is due to the formation of lead sulphate in the mixture.

Lime Resistant Paint

Complete protection against corrosion by hot lime-water and acetylene residues is obtained by a paint containing 16 per cent chlorinated rubber, 44 per cent xylene, 35 per cent lithopone, and 5 per cent tritolylphosphate.

Luminous Paint Swiss Patent 172,076

Sandarac	36 g.
Rosin	18 g.
Paraffin	4 g.
Alcohol	35 g.
Petroleum Ether	10 g.
Tricresyl Phosphate	1 g.
Benzoin, Gum	2 g.

Mix with gentle warming until dissolved. Dehydrate with quick-lime and filter.

65 grams of above are mixed with: Strontium Sulphide 35 g.

Mildew Preventatives for Paint

The addition of any of the following per 600 pounds of paint is advisable:

Mercuric Chloride	1 lb.
Sodium Silico Fluoride	6 lb.
Ammoniated Mercury	2 lb.

Non-Caking Pigments

Pigments are prevented from caking and are more readily dispersed in either oil or water if they are suspended in a dilute dispersion in water of diglycol stearate or glyceryl monostearate and then dried. A film of waxy material is formed around each pigment particle. This film is both oil soluble and water dispersible.

Marble-Effect Dipping Paint

Beautiful, marble-like effects are obtained by dipping objects into many-colored paints floating upon the surface

In nn

of water. In order to float on water, the paints used have to weigh less than 8.33 pounds per gallon. Assuming that a varnish is used which weighs 7 pounds per gallon, the following table gives the number of pounds of pigment which, when ground into 1 gallon of varnish, will yield a paint of sufficiently low weight to float on water, and have good hiding.

Chrome Yellow	1.25
Chrome Green	1.00
Prussian Blue	0.50
Para Red	0.50
Aluminum Bronze Powder	1.50
Gold Bronze Powder	1.50
Carbon Black (High Strength)	0.50

The procedure is important. Select a container which is wide enough and deep enough to hold the largest object to be dipped. Fill the container with water at room temperature. By means of a rod or dropper place a few drops of a colored paint here and there on the surface of the water. Near these drops or upon them place drops of a contrasting colored paint. Three, four or even five different colors may be used, but an excess of paint should be avoided. The colors will spread about, mingling with each other. They may also be blown gently. Hold the object to be decorated in such fashion that the entire outside surface is exposed. Immerse it slowly into the colors and into the water, turning it a bit at the same time. Blow the remaining colors aside in order to withdraw the object without having it traverse the colors The designs produced in this manner will always be different from each other, and are almost impossible to reproduce by hand painting.

Oiticica Oil Emulsion Paint U. S. Patent 1,998,845

Oiticica Oil	120 oz.
Lead Oxide	6 oz.
Manganese Dioxide	2 oz.
Heat to 250° C and then	reduce to

200° C. and add

Potassium Silicate 13 oz.
Milk of Lime 16 oz.
Water sufficient

Agitate violently until cool.

Paint Perfume

Vanillin is dissolved in turpentine or linseed oil. One part of vanillin is used to 2000 parts of paint to cover objectionable odors.

Plastic Paints

Zinc White or Lithopone	18.15	oz.
Water	7.5	oz.
Hide Glue	0.68	oz.
Linseed Oil, Pale Boiled	3.8	OZ.
Rosin (WW or WG)	3.6	oz.
Benzol	3.8	oz.
Zinc Sulphate	0.12 .	oz.

If a hard dry product is wished, add gypsum. Treat with water until pasty.

Synthetic Resin Enamel Paints Formula No. 1

Zinc Oxide (White Seal)	400	lb.
Thin Stand Oil	180	lb.
Turpentine	100	lb.

Pug well and grind four times, then add:

China Wood Oil Varnish, con-		
taining 25 per cent Syn-		
thetic Resin, equivalent to	88	lb.
Thick Stand Oil		lb.
Turpentine	64	lb.
Cobalt Lingleage (Liquid)	20	lh

This enamel dries in from 15 to 18 hours

No. 2

Titanium Oxide	300 lb.
Zinc Oxide	300 lb.
Thin Stand Oil	180 lb.
Synthetic Varnish	250 lb.
White Spirit	100 lb.
Cobalt Linoleate	10 lb.

No. 3

Zinc Oxide	200	11.
Zine Oxide	300	
Titanium Oxide	300	lb.
Thin Stand Oil	280	lb.
Synthetic Varnish	150	lb.
White Spirit	100	lb.

Synthetic Resin for Paints Canadian Patent 348,347

Castor oil 500 and drying oils 500 parts by weight are mixed and distilled until the residue of polymeric esters is approximately 85% of the original mixture. The retort is cooled below 290° and 800 parts of glycerol is gradually introduced. The mixture is heated for a short time well above the boiling point of water but below the boiling point of glycerol, and then 1200 parts of phthalic anhydride is gradually added, the temperature being maintained about midway between the boiling point of phthalic anhydride and that of water. When the mixture is clear and homogeneous it is run into cooling pans or into mixing tanks to be thinned with solvents.

Tar and Asphalt Paints Formula No. 1

Pine tar 120 l., rubber (small pieces) 1300 g., gutta-percha (small pieces) 1600 g., shellac 2700 g., copal varnish 4.5 l. When the varnish has been incorporated 45 l. of linseed oil heated separately to nearly the same temperature are added slowly.

No. 2 .

Asphalt 40 g., fossil resin 10 g., heat-thickened linseed oil 8 g., liquid driers 20 g., turpentine 60-70 g.

Paint for Marking Wood Boxes, Barrels, etc. Formula No. 1

Gum Arabic	10	g.
Soda Ash		g.
Glycerin	1	g.
Water	40	g.

Lampblack or pigment, as much as needed.

No. 2

Waterproof:

Silica

" accept com .	
Shellac, Ruby	60 g.
Borax	60 g.
Water	750 g.
Dissolve boiling, and add:	Ŭ
Gum Arabic .	60 g,
Pigment or Lampblack, as	
needed.	

Cement Water Paint German Patent 575,895

40 kg.

Pyrolusite Whiting			5 kg. 40 kg.
Cement			15 kg.
Grind very finely following solution:	and	mix	into the
Casein			50 kg.
Borax			30 kg.
Water			150 kg.
Rosin Emulsion			20 kg

Wool Fat Emulsion Paints German Patent 612,715

Ammonium salts of high molecular fatty acids derived from drying or semi-drying oils have been claimed to be exceptionally valuable emulsifying agents for paint compositions incorporating both wool fat and non-water-soluble ingredients, such as resins and drying oils. Not only are the resulting coatings far more water-resistant than those of ordinary wool fat coatings, but the employment of an aqueous medium obviates

some of the drawbacks of solution in organic solvents. The process can be illustrated with reference to an emulsion of crude wool fat, refined tung oil and rosin, which are melted up in the respective proportions of 360:40:250, the melt being incorporated with 43 parts of ammonium solution, 100 parts alcohol and 1207 parts water and the resulting emulsion agitated till cold. The product at this stage, a viscous, yellowish-white emulsion, may be directly employed as a paint. An example of a quick-drying paint comprises 1000 parts emulsion, 80 parts chrome oxide, 150 parts titanium white and 15 parts of a 33 per cent solution of a cobalt-leadmanganese drier. Such a paint is stated to reach surface dryness within two hours after brushing on any type of surface, and admirably resists the action of a condensed steam-laden atmosphere.

Specialty Paints

French Patents 44,177 and 756,535

Under-Water Paint:

Water	500 kg.
Tar	300 kg.
Caoutchouc Solution	200 kg.
Rosin	200 kg.
Benzene	100 kg.
Alum	TOO Kg.
	2 kg.
"Very Brilliant" Paint:	
Alum	12 g.
Aluminum Bronze	5 g.
Salt	30 g.
Sugar	5 g.
"Fatty" Lime	50 g.
Water	400 g.
Oil	400 g.
Rosin	400 g.
Benzene	200 g.
Mica Powder	150 g.
	20 g.
Milk Whey	100 g.
Caoutehoue Solution	200 g.
Liquid Drier	150 g.
Pigments	10-15 g.

Paint and Varnish Remover

Formula No. 1		
Amyl Acetate	15	lb.
Acetone	14	lb.
Benzol	īī	lb.
Methanol	12	lb.
Paraffin Wax	21/2	
No. 2		
Whiting	21	lb.
Acetone	21	lb.
Denatured Alcohol	21	lb.
Benzol	23	1h

114 lb.

Paraffin Wax

No. 3

A low priced and effective remover may be made up as follows:

Ethyl Acetate	30	oz.
Benzol	40	oz.
Methanol	27.5	OZ.
Paraffin Wax	2	oz.
Methyl Salicylate	0.5	oz.

The paraffin is melted and poured into the benzol. The other solvents are mixed and then the benzol wax solution added to same while mixing vigorously.

Removing Plastic Paint

Mix one pound sal soda and two pounds hydrated lime and one-fourth of a pound of table salt. Add enough water to this mixture to produce a fairly heavy paste. Apply the paste with a fiber brush, and leave it on until the old material is softened, when it may be scraped off. If the paste material should become nearly dry before the old material is soft enough to be easily scraped off, apply the paste material again, but always be sure you do not get this caustic paste on the woodwork or floors, as it would injure them. When all the old material has been scraped off, wash the surface and rinse it until it is perfectly clean, and allow it to become dry before applying the first coat of paint.

Finish Remover U. S. Patent 1,974,744

Acetone	35 oz
Ethyl Acetate	15 oz.
Denatured Alcohol	10 oz.
Benzol	10 oz.
Oxidized Pine Oil	10 oz.
Diethyl Phthalate	20 oz.
Diethyl Phthalate Cellulose Acetate	4 oz.

Varnish Remover, Liquid

Methanol	30	gal.
Phenol (90%)	5	gal.
Light Coal Tar Oil		gal.

Varnish Remover, Paste

Crude Vaseline	50 gal.
Phenol (90%)	45 gal.
Fusel Oil	20 gal.
Wood Flour	80 lb.

Shellac Finish

Shellac			250	g.
Dragon's	Blood	*	50	
Alcohol			750	

Mix until dissolved, while warming on water bath.

Copal (Po	wdere	d a	nd	Ex-	
posed to	Air	for	a	Few	
Weeks)					60 g.
Alcohol					250° c

Dissolve by mixing on water bath and then add:

Chalk, Precipitated 180 g. Then mix with first solution.

Flat	Indoor	${\bf Shellac}$	Lacquer	
Copal			131/2	oz.
Alcohol			131/2	oz.
Shellac T	'.N.		7	ΟZ.
Alcohol			18	OZ.
Bone Oil			- 3	oz.

Finishing	Shellac	Lacquer	
Shellac, White	Refined	100	oz.
Alcohol		125	oz.
Butyl Alcohol		4	OZ.
Bone Oil		1	oz.

Danahina Diniahina Challes Ta

Brusning Finishing Shellac	Lacq	uer
Copal	21/2	OZ.
Alcohol	21/2	oz.
Sandarac	1/2	oz.
Alcohol	1	oz.
Shellac, T.N.	2.2	oz.
Alcohol	3.3	oz.
Acaroid Red, Alcoholic		
(1:1)	1	oz.
Acaroid Yellow, Alcoholic		
(1:1)	1/2	oz.
Butyl Alcohol	1/2	oz.
Castor Oil	1/2	oz.
Bone Oil	1/4	oz.

Shellac Floor Finish

Shellac Floor Fil	11911
Shellac, Orange	280 g.
Linseed Oil Varnish, Pal	le 80 g.
Ochre, Pale or Dark	50 g.
Alcohol	- 1 I.
Stir altogether, let stand	over night.

Floor Refreshener

5 lb. Shellac ''Cut'' Denatured Alcohol	1/4 3/4	gal. gal.
		_

This mixture is applied with a mop. The alcohol cleans and at the same time there is left a thin film of shellac which adds lustre to the floor.

Shellac Polish

Lac, Button	18	oz.
Alcohol	72	oz.
Shellac, T.N.	9	oz.
Sandarac	4	OZ.
Benzoin, Gum	4	OZ.
Turpentine, Venice	5	oz.

Water Shellacs	
1. Bleached "Pig-Tail" Sh	ellac
Water	645 g.
Borax	55 g.
"Pig-tail" Shellac, Ground,	
20% Water	300 g.
2. Bleached Shellac Power	ler
Water	705 g.
Borax	55 g.
Shellac Powder, Dry	240 g.
Ruby and Orange Shel	lac
Water	700 g.
Borax	50 g.
Ruby or Orange Shellac	
(Free of Rosin and Wax)	250 g.
Solution in above formula is	hastened

Water Resistant Shellac

by warming and stirring.

Add 2-3% of urea or thiourea to solution of shellac in alcohol.

Bleaching Shellac

Lac may be bleached by dissolving it in 2.5% sodium carbonate solution at 60-70° and, after filtration and cooling to air temperature, adding a solution prepared by passing chlorine into a solution containing 12.5% of caustic soda and 2.5% of sodium carbonate. latter should contain 6-8% of available chlorine and, if of pH 10-10.5, does not require storing in a cool place. The amount of such a solution necessary for bleaching indicates a chlorine requirement of 10-14% on the weight of lac, and a yield of 93-95% is obtained. The bleached lac may be recovered by the slow addition, with stirring, of 1:20 sulphuric acid, the precipitate being then collected, washed, and dried in vacuo over sulphuric acid. The product is freely soluble in cold 97% alcohol, and the solubility does not alter on prolonged storage in air. The bleached material contains 2.3-3.1% of moisture, 0.98-3.52% chlorine and has a saponification value 236.0-256.7, acid value 70-68-83-52, and iodine value 3.9-5.0.

Substitute Shellac Solutions

The substitutes for shellac solutions are of three types:

- 1. Substitute for wax-free shellac solution.
- 2. Substitute for white shellac solution.
- 3. Substitute for orange shellac solution.

The base for all three is the same, namely a solution of a cheaper alcoholsoluble resin in completely denatured alcohol. At the present time a soft Manila gum is used, and a 6-lb. cut represents the maximum concentration normally made. To prevent loss by evaporation, as well as to avoid the hazard of volatile alcohol vapors, a closed tumbler is used, in which is placed one gallon of alcohol for every six pounds of the Manila gum. When solution is complete, the tumbler is emptied and the solution allowed to settle. The clear supernatant solution represents a substitute for waxfree shellac solution.

White and orange shellac solutions contain a cloud of suspended wax which is inherent in the material and insoluble in alcohol. To duplicate the waxy appearance a preparation of carnauba wax may be employed. A quick and safe method of preparing the wax is as fol-

Imitation Shellac "Cloud"

Dissolve 5 lb. of carnauba wax in onehalf gallon of blown easter oil. Since carnauba wax melts at 84-86° C., a steam-jacketed kettle may be used. If a direct fire is used, the flame must be extinguished before proceeding further with the formula. Add slowly and with constant stirring one-half gallon of turpentine, followed by one-half gallon of denatured alcohol. A soft yellowishwhite paste will form. This paste, added to a solution of 95 lb. of Manila gum in 15 gal. of alcohol, represents a 6-lb. cut in which the wax constitutes 5% of the total solids. Less paste may be used, but not more. The castor oil serves not only as a solvent for the wax, but also as a plasticizer.

The waxed product is a substitute for white shellac. It may be colored by means of an orange alcohol-soluble aniline dye, thus forming a substitute for

orange shellac.

Shellac Substitute U. S. Patent 1.942,413

Batu (Galla-Galla) Gum 18-20 oz. Rosin 10-20 oz. Heat to 260° C. Add: Calcium Oxide

1-4 oz.

Heat to 320° C. and stir till dissolved. Cool and "cut" with varnish solvents to give a shellac substitute solution.

Oiticica Varnish

An oiticica oil varnish cooked under the same conditions as a similar tung oil varnish is lower in viscosity, which is an advantage. If the temperature is taken over 250° C. frothing occurs and this has to be carefully watched.

By blowing oiticica oil for 30 minutes at 220° C. a thick light-colored oil is formed which will be comparable with blown linseed oil. Oiticica oil varnishes have a less characteristic odor and are less noticeable in closed spaces.

To establish the technical value of oiticica oil, tests have been made with varnishes with a natural or artificial resin base and mixtures on the one hand of tung oil and linseed oil, and on the other of oiticica oil and tung oil, the latter being in the ratio of one part to two respectively. Heating is done at 315° C. and maintained until the mixture has the correct body.

Ester Gum Varnishes Formula No. 1

Ester Gum	100 lb.
Tung Oil	198 lb.
Linseed Oil Heated for	
2 Hours	36 lb.
Solvent Naphtha	84 lb.
White Spirit	250 lb.

Driers are added in the proportion of 0.5% lead and 0.035% cobalt. This gives a varnish which becomes tacky in 45 minutes and dries in about 3 hours. The film is resistant to cold and boiling water. The film is not resistant to combustion gases. The Gardner-Holt viscosity is D and the color 11.

No. 2

Ester Gum	100 lb.
Oiticica Oil	156 lb.
Tung Oil	78 lb.
Solvent Naphtha	84 lb.
White Spirit	250 lb.

Driers are used in the same proportion, i.e., 0.5% lead and 0.035% cobalt. This varnish becomes tacky in 2 hours and perfectly dry in about 8 hours. The film is resistant to cold water but not to boiling water when it whitens but becomes transparent again.

Pharmaceutical Cellulose Varnish French Patent 777,999

A varnish containing, e.g., benzylcellulose 5–18 g., benzine 18–40 g., toluene or xylene 25–45 g. and butyl acetate 20–35 g., or benzylcellulose 2–12 g., benzine 50–80 g., and ether 25–80 g., used for pharmaceutical or toilet purposes, is contained in a collapsible tube and used as required.

Electrically Conducting Varnish Formula No. 1

Aluminum Bronze Powder	240 g.
Synthetic Resin Varnish	1 l.
No. 2	

Copper	Bronze	Powder	120 1	g.
Lacque	r		1	1.

Cold "Cut" Synthetic F	tesin Vari	nish
Rezyl No. 14	10	g.
Methanol	50	čc.
Toluol	5 0	cc.

Allow to stand over night and then stir.

Leather Roller Varnish

Venetian Red	4	lb.
Ground Blue	5	lb.
Vinegar		pt.
Glycerin	75	cc.
Glucose	150	
Oil of Cloves		cc.
Methyl Salicylate	25	cc.

Mu Oil Varnish

Mu Oil			200	oz.
Modified	Phenolic	Resin	100	oz.

Heat with stirring to 570° F.; keep at this temperature for 6 minutes; cool to 350° F. and dilute with 250 oz. petroleum spirits and add 5½ oz. lead naphthenate and 0.1 oz. cobalt naphthenate.

Mopping and Wiping Varnish

Because varnishes of this type leave a very thin film, it is essential that they be made of tough and durable ingredients. The average floor or furniture varnish, if thinned to wiping consistency, is unsuitable. A high grade product consists of a blend of 3 or 4 pints of the following varnish a with 1 pint of varnish b.

a. Bakelite XR-4070 100 lb.
 China Wood Oil 16 gal.
 Body for 1 hour at 450° F. Reduce with 20 gal. mineral spirits, 5 gal

xylol, 3 gal. dipentene, 2 gal. high boiling hydrogenated naphtha and 3 gal. gum spirits of turpentine.

b. Substitute Bakelite BR-\$20 in place of XR-4070 in Formula a, and body with the wood oil at 400° instead of 450°.

Driers are unnecessary.

High Gloss Transparent Printing Varnish

British Patent 426,753

Ester Gum	120	oz.
Tung Oil	40	OZ.
Linseed Oil (Half Boiled)	40	oz.
Mineral Spirits	6	oz.
Cobalt Linoleate	5	oz.

The above may be colored with a suitable amount of rhodamine base dissolved in olein, Berlin blue, alizarin madder take, milori blue or Sudan yellow.

Wrinkle Finish Varnish U. S. Patent 1,934,034

Tung Oil	100 oz.
Rosin	5-10 oz.

Heat for 2 to 8 hours at 177-230° C. Cool and dissolve in an equal weight of high-flash naphtha.

Limed Rosin

The apparatus and procedure vary somewhat, but the following is usual practice: One hundred and twenty-five pounds of resin are melted in a cylindrical flat-bottomed copper vessel 36 inches in height and from 30 to 36 inches in diameter. The vessel has a loose cover provided with a small chimney and an opening for the stirring rod. It is mounted on an iron truck, the platform of which is about 2 inches from the floor. The truck is then wheeled to a position under a chimney and over a furnace, which is located beneath the floor. The resin completely melts in about a half hour. It is at this point that the use of lime enters.

Lime is added, gradually, to the melted resin with the temperature at about 350° F. Theoretically, about 13.6 pounds of hydrated lime would be required, but it is inadvisable to completely neutralize the resin. In actual practice 8 to 10 pounds of hydrated lime are used. This reduces the acidity of the resin from about 160 to 65. After stirring and heating for a short while, the treatment with lime is completed.

Wood Filler

Shellac (if for Tran	
Wood Filler Use B	Bleachea
Shellac)	4 lb.
Methylated Spirits	1 gal.
Barytes	20 lb.
Silica	10 lb.
Raw Linseed Oil	1/4 gal.

Dissolve the shellae in the methylated spirits and add the linseed oil. Mix the barytes and silica together dry, and stir into the shellae varnish. Grind to a smooth paste and adjust the consistency with additional barytes and silica mixture or shellae varnish. Store in airtight containers.

Filler-Undercoat for Shellac

Mixing powdered boracic acid, 5 g., with each ounce of shellac to be used as an undercoat on wood cuuses the shellac to dry very hard so that it serves as a filler as well as an undercoat.

Porous Wood Sealer

One hundred thirty-five pounds of 400-mesh Silica, 65 lb. Bentonite, 10 gal. of Congo Copal Varnish, 2½ gal. Pontianak Gum Varnish, 2½ gal. Nevindene Solution, 10 gal. Light Naphtha, 5 gal. Lacquer Thinner, ½ gal. Concentrated Cobalt Drier Nevindene solution is (basis) 6 lb. of Nevindene resin cut cold in 1 gal. of mineral spirits.

The protective covering should be a coat of aluminum paint and advisedly two coats of regular oil-type house paint. The Aluminum Primer recommended is: 72½ gal. of an 80-gal. Tung/Ester Varnish, 10 gal. Boiled Linseed Oil, 7 gal. Mineral Spirits, ½ gal. Lead-Mauganese Concentrated Drier, 125 lb. Paste Aluminum (or powder).

Non-Penetrating Plaster Sealer Pigment 45 lb. Vehicle 55 lb. Pigment: Titanium-Calcium Pigment 62 lb. 37 lb. Metronite Aluminum Stearate 1 lb. Vehicle: Bodied Linseed Oil 50 lb. Mineral Spirits 45 lb. Liquid Drier 5 lb.

Wood Filler for Ground Polishing German Patent 607,521

	,
Shellac Wax	10 oz.
Carnauba Wax	5 oz.

½ gal.

Water

Pumice Meal	100 oz.
Sandarac	100 oz.
Blown Castor Oil	10 oz.
Melt together until unif	orm and pow-
der after cooling.	_

American Walnut Graining Color Ivory Black 2 oz. Van Dyke Brown 4 oz. Burnt Umber 2 oz. Bolted Whiting 1 oz.

Imitating Old Copper Finish After application of priming coat use lb. White Lead Chrome Yellow, Medium 12oz. 1½ lb. Venetian Red Burnt Umber 4 oz. Linseed Oil 41/4 lb. 41/4 lb. Turpentine to suit

After applying above paint, allow to dry and use a coating of copper bronze powder thinned with equal parts of spar varnish and turpentine. When this coat is dry apply a glaze made from chrome green, medium, thinned with equal parts of raw linseed oil and turpentine plus a small amount of drier. While the glaze is still damp wipe it here and there to produce a mottled effect.

Liquid Oil Graining Color
Raw Linseed Oil % gal.
Turpentine % gal.
Drier, Liquid ½ pt.
Beeswax, Yellow (Shavings) % oz.
Warm together and mix until clear.

Wood Stain U. S. Patent 1,977,345

Dye, Water Soluble 4 oz.
Diethylene Glycol Ethyl Ether 5 oz.
Alcohol 80 oz.
Ethylene Glycol Methyl Ether 15 oz.

Wood Stain U. S. Patent 2,000,121

Diethylene Glycol Monoethyl Ether 1 oz. Methyl Alcohol 9 oz. Toluol 6 oz.

This composition may be utilized with from 2 to 2½ oz. of the particular dye to 1 gal. of the composite solvent. The amount of dye utilized depends on the particular dye itself and its degree of

concentration, and the depth of color required in the particular stain. Further, the strength of the dye stain may be varied by the amount of diethylene glycol mono-ethyl ether utilized.

In making up these compositions, the aniline dye or stain, such as the nigrosines, may be allowed to stand with the diethylene glycol mono-ethyl ether until the dye dissolves, after which the other ingredients may be added.

Coloring of Light Wood to Imitate Ebony

A vacuum process is essential for good impregnation of wood with coloring substances. Aqueous solutions are preferable where possible on grounds of low price, high vapor pressure (which assists impregnation) etc. Woods for this ebonizing process, in order of suitability are: apple, pear, hazel, maple, beech and birch. The following are recipes for ebonizing:

Formula No. 1

Gall-nut solution containing a few drops of ammonium vanadate solution.

No. 2

3.60 kilograms aniline hydrochloride, 1.80 kilograms potassium chlorate, 40 liters water, 0.250 liter hydrochloric acid, 4.20 grams ammonium vanadate.

No. 3

Four kilograms carbon black, 18 liters shellae Japan lacquer, 18 liters turpentine.

No. 4

1,200 kilograms carnauba wax, 3 kilograms ceresin, 30 grams oil-soluble nigrosine, 10 liters turpentine.

Auto Top Dressing

Orange Shellac 4 oz.
Denatured Alcohol 1 gal.
Castor Oil 4/2 oz.

If a black finish is desired add nigrosine to give the desired color.

 $\begin{array}{cccc} \text{Butter Taint Prevention Coating} \\ \text{Tubs are coated with following:} \\ \text{Prime Lactic Casein} & 50 & \text{oz.} \\ \text{Borax} & 7.5 & \text{oz.} \\ \text{Water} & 300 & \text{oz.} \\ \text{Stir and warm gently until smooth.} \\ \end{array}$

Candy Glazes Formula No. 1

Sandarac 125 g. Benzoin, Sumatra 125 g.

Turpentine, Venice Alcohol No. 2	10 g. 740 g.
Benzoin, Sumatra Balsam, Peru Alcohol No. 3	200 g. 5 g. 800 g.
Benzoin, Sumatra Shellac, Refined Vanillin Alcohol	150 g. 50 g. 1 g. 800 g.

Protective Food Coating French Patent 780,762

Lactic Casein Borax or Sodium Phosphate Sodium Bicarbonate Glycerin Distilled Water Gelatin	100 16 32 34 820 8	000 00 00
Gelatin	8	g.

This may also be applied to aluminum or tin foil for use on foods.

Protective Coatings for Sausages, etc.

	Formula No. 1	
Gelatin Salt		5 g. 2 g. 1 g.
Saltpeter	No. 2	1 g.
Gelatin	2100	5 g. 1 g.
Glycerin	No. 3	1 g.
Pentosan	n Resin	3 g.
Gelatin	No. 4	1 g.
	110. 1	

Aqueous Solution of Stahr, or Agar-Agar, or Gelatin, ½-2% Formic Acid.

No.	5
Talle	ow

	710.	0			
Alum				1	g.
Olive Oil				1	cc.
Shellac				16	g.
Alcohol				65	cc.
	TAT o	77			

Paraffin		35	g
Colophony		62.8	
Whiting		2.2	
	No. 8		0

Linseed Oil	60	g.
Colophony, Shellac, Glycerin,		
or Wax	40	g.
No. 9		

Glue, Gelatin or Isinglass, boiled in a little vinegar.

Laboratory Table Top Stain Solution No. 1

Potassium Permanganate	20 g.
Copper Sulphate	20 g.
Water	1 l.

Heat to about 60-70° C. and apply to clean desk top, and follow immediately with solution No. 2.

Solution No. 2

Hydrochloric Acid	
(sp. gr. 1.2)	150 cc.
Aniline	150 ec.
Water	700 cc.

Heat to about 60-70° C. and apply over No. 1.

When the desk top is dry it may be rubbed with linseed oil in the usual manner.

Red Stamp Pad Ink

Fuchsin	1	oz.
Glycerin	32	OZ.
"Lysol"	1/8	OZ.
Acetic Acid	1	oz.
Denatured Alcohol	1	$0Z_{\bullet}$
Water	1	oz,

Protector for Polished Surfaces French Patent 778,389

Water	150	cc.
Linseed Oil	200	cc.
Alcohol	450	cc.
Sulphuric Acid		cc.
Shellac	30	g.

Coating for Old-Painted Surfaces Swiss Patent 173,070

	•
Trichlorethylene	25 cc.
Polishing Lacquer	25 cc.
Benzoline	25-30 сс.
Lithopone, as Pigment	optional

Preparation of Oil Pastes from Pigment-Water Pulp

The addition of linseed oil of acid value about 10 will cause the separation of water from a pulp of white lead-inwater. Agitation and friction are necessary in order to insure contact of the oil with the pigment and in order to express the maximum amount of water.

With other pigments, particularly those whose affinity for oil is less striking than that of white lead, transfer from the water phase to the oil phase may be

accomplished by one or more of the following means:

High acid linseed oil.
 Polymerized linseed oil.

3. Linseed or China wood fatty acids.

4. Addition of various chemical agents.

As an example of method 4 (Patented), 15.5 parts of linseed oil (acid value 7) or of other drying oil (acid value greater than 4) are added gradually at 82-88° C., with vigorous agitation, to a suspension of 100 parts of lithopone in 200 parts of water which also contains tri-sodium phosphate or other alkaline saponifying agent. The water separates in the upper layer after 10 to 30 minutes' further agitation.

Strong Lead Oil for Black Paints

Varnish linseed oil is heated with continual stirring until at the end of an hour the temperature reaches 570° F. (=300° C.) and is held at this temperature for a further 3 to 4 hours, when the heat is closed down. Finely powdered white lead is then added slowly on a falling temperature, commencing at about 525° F. (=274° C.), in the proportion of 4½ lb. of white lead to every 100 lb. of oil, about 2 hours being occupied in adding the white lead. So far, it will be observed, the process will have occupied practically one working day. On the following day the oil is heated up again, care being taken to avoid local heating in the early stages until the whole mass Heat is then inbecomes quite fluid. creased until a temperature of 535° to 545° F. (=280° to 285° C.) is reached, at which the oil is held until the body required is attained. The purposes for which oil of this type is used demand as a rule that the product when cooked shall "string" very strongly when tested Gums or blacks with which it may be cooked afterwards are usually expected to "pill" between the finger and thumb.

Flatting Oil

Linseed Oil		15 lb.
Solvent Naphtha Turpentine	or	85 lb.
Drier		to suit

Add to the following lead paste in proportions of 2½ gal. above oil to 100 lb. lead paste:

White Lead 92 lb. Linseed Oil 8 lb.

Black Iron Oxide Pigment Austrian Patent 141,130

Ferrous Sulphate Water	240 lb. 720 lb.
Boil the above and while	boiling add:
Potassium Chlorate	12 lb.
and then add: Sodium Carbonate	115 lb.
in Water	230 lb.

Various shades are obtained by varying composition of first solution, nature and amount of oxidizing agent and other reaction conditions.

Carmine Lake Pigment

Powder the best silver-gray cochineal as finely as possible, and boil it for three hours in water. Filter the hot solution quickly through a thick linen cloth. Boil up the filtrate again, and add the substances needed to form the lake. Many such substances may be used, but only two can be thoroughly depended upon, and they should both be used together. These two are alum and tin salt, and if necessary, warmth may be given to the color by the cautious addition, drop by drop, of hydrochloric acid. The alum must be absolutely free from iron, or it will be impossible to get more than a very unsatisfactory product. The best proportions are:

 Cochineal
 20 lb.

 Water
 500 lb.

 Alum (Iron Free)
 2 lb.

 Tin Salt
 2 lb.

The alum and tin salt are added at the boil, which is kept up till everything is dissolved. The clear solution is then exposed in shallow dishes covered with sheets of glass for several weeks in a very bright sunny place. By this time the dark-red liquor will have lost nearly all its color, and the carmine will have been deposited in the solid form, partly on the dish and partly on the surface of the liquid. It is separated by filtration, and carefully dried with blotting-paper. To get a fine and warm red it is absolutely indispensable that the dishes should get plenty of sun, so that the manufacture is impossible in any but the most favorable weather.

To get absolutely pure carmine, the product already described is dissolved in caustic ammonia. The solution is filtered, and the carmine is reprecipitated with acetic acid.

Satin White Pigment

Ninety pounds of quicklime are slaked in 27 gal. of boiling water. To this mixture 130 lb. of finely divided (260 mesh) aluminum sulphate are added quickly, and the mass is heated until it becomes almost solid. Thirty gallons of water are then added and the mixture agitated thoroughly. The last trace of any visible yellow color is neutralized by the addition of indanthrene blue in the form of a solution of 2.5 lb. of the commercial paste in 6 gal. of water. A very small amount of this solution is required if a good grade of lime and sulphate are used. The satin white is then filtered and dried.

Reflecting of Light by Colors

	Reflection
	Factor
Color of Paint	Per Cen
White (Gloss)	84
White (Flat)	82
White (Eggshell)	81
Ivory White	79
Cream	74
Aluminum (Made with Paste	
Ivory Tan	67
Light Green	62
Light Gray	59
Buff	55
Light Blue	52
Medium Green	49
Tan	48
Medium Blue	43
French Gray	32

Printing in Several Colors British Patent 426,753

High-gloss color-printing is effected by printing the picture in black or other color in the usual way and over-printing the picture entirely or partly with a transparent colored gloss overprint varnish. The varnish may be colored with oil-soluble coloring matter or with highly glazing insoluble pigments or with both. In the last case, autotype prints having a double tone effect may be produced, the soluble color spreading out around each of the dots of the picture. The first print may be made with a normal black art printing ink. The varnish consists of clear resin ester 120, china wood oil 40, slightly boiled linseed oil 40, petroleum 6 and cobalt linoleate 6 parts. To 12 parts of varnish may be added 2 of rhodamine base in 2 of olein, 1 of Berlin blue or 1 of alizarin madder lake (I). Double tone effects may be produced by over-printing with a mixture of varnish 25, rhodamine base 0.5 in olein 0.5, and milori blue 1 parts, or with varnish 25, Sudan yellow 0.5 and (I) 1 part.

Dissolving Amber

The amber is powdered and heated under a reflux condenser with butyl alcohol containing a little hydrochloric acid for 6 to 8 hours.

Dustless Carbon Black Formula No. 1

Carbon Black	200	g.
Sapropélite Tar	24	g.
Water	50	čc.

Form pellets or briquettes and dry at 105° C. for 3 hours.

No. 2 Carbon Black 200 g. 100 cc. Dextrin Solution (5%) Treat as above.

Colloidal Preservative U. S. Patent 1,937,813

A transparent, solvent-resistant, antiseptic, colloidal mass is produced by condensing the gases evolved when gelatin 3 lb. or glue, etc., is heated with wood creosote 4 lb. at 160-250° C. for 2 hours.

Coloring	Meerschaum	Pipe	Boy	vls
Beeswax			50	oz.
Olive Oil			50	OZ.
Triethand	lamina		7.5	07

The Meerschaum pipes are immersed in the above which is slowly heated to boiling and maintained at this temperature for 15 to 30 minutes. Pipes so treated will color very rapidly.

Blue Sheep Marking Pencil Soapstone 28 lb. Fine Gypsum 21 lb. Chinese Blue 2 lb.

White Soap Powder 10 lb. Mix all ingredients well together and make up with thin glue water into a stiff paste. They are then shaped like a

thick pencil and dried.

Brewer's Glaze

Orange Shellac	25	oz.
Manila Copal	12	OZ.
Acaroid Resin, Yellow or		
Red	8	oz.
Linoleic Acid	0.5	oz.
Alcohol	54.5	oz.

Rubbing Compound (For Paint, Lacquers, etc.)

(x or x arms, xacquers,		
1. Carnauba Wax	42	lb.
2. Beeswax	18	lb.
3. Ceresin	18	lb.
4. Varnolene	3	gal.
5. Water		gal.
6. Triethanolamine		oz.
7. Stearic Acid	2	lb.
8. Tripoli	24	lb.
9. Pumice	15	lb.

Melt 1, 2, 3, 7 with 4. Heat 5 and 6 to 90° C., add to wax mixture and stir till emulsified. Then add 8 and 9 and stir till cool.

Peeled Wood Wall Paper U. S. Patent 1,945,686

The veneer is cut into strips of definite width which are dried, steeped in solution (1), dried, steeped in solution (2), dried, and finally backed with any kind of fibrous fabric. (1) comprises cellulose acetate 15, 14% solution of chrome alum 10, and water 70 oz., and (2) 25% glycerin 30, gelatin 25, and water 45 oz.

Double Strength Lead-Manganese Liquid Drier

Lead-Manganese Uversol		
No. 303	200	lb.
Bodied Linseed Oil	73.5	lb.
Pine Oil	9.0	
Turpentine	60.0	lb.
Pine Tar Oil	3.0	lb.
Mineral Spirits	254.5	lb.
Yield 75 gal.		

This drier is double the strength of the preceding, containing 1.0% manganese and 11.0% lead as metals.

Procedure: Melt the drier quickly with the linseed oil at a temperature not exceeding 275° or 300° F. Remove from fire and reduce with the solvents.

Lead-Manganese Drier

Lead-Manganese	Uversol		
No. 303		100	lb.
Mineral Spirits		500	lb.
Viold 85 gel			

This drier has an acid value =0. It contains 0.5% manganese and 5.5% lead as metals. One part of drier to twenty parts of oil will give a metallic content of 0.025% manganese and 0.275% lead.

COSMETICS AND DRUGS

AND DROGS	
Medical Bathing Sa Carlsbad Well	
Sodium Sulphate Potassium Sulphate Sodium Chloride Sodium Bicarbonate	44 g. 2 g. 18 g. 36 g.
Notice production	
Friedrichshall	
	37.7 g. 0.3 g.
Potassium Chloride	5 g.
	19 g. 37 g.
Calcium Sulphate,	J
Precipitated	1 g.
7.1.1.1	
	6 g.
Magnesium Chloride	72 g.
	0.15 g. 14 g.
Sodium Bromide	0.85 g.
Magnesium Sulphate	7 g.
Krongnogh	
	63 g.
Potassium Chloride	75 g.
Magnesium Chloride	750 g. 110 g.
Sodium Bromide	2 g.
-	
Hallein Well	
	69.3 g. 27 g.
Sodium Bromide	0.42 g.
	10 g.
Sodium Sulphate	2.28 g.
Water Super distance from the control of the part of the control o	
Vichy	3 - 1 m - 1
	0.01 g. 0.05 g.
Manganese Šulphate	0.01 g.
	1.73 g.
Magnesium Sulphate	2.6 g.
Calcium Chloride Sodium Bicarbonate	6.0 g. 83.4 g.
	Medical Bathing Sa Carlsbad Well Sodium Sulphate Potassium Sulphate Sodium Chloride Sodium Bicarbonate Friedrichshall Sodium Chloride Sodium Bromide Potassium Chloride Calcium Sulphate, Precipitated Reichenhall Potassium Chloride Magnesium Chloride Lithium Chloride Sodium Bromide Sodium Bromide Sodium Bromide Sodium Chloride Sodium Chloride Sodium Bromide Magnesium Chloride Sodium Bromide Calcium Chloride Sodium Bromide Kreuznach Sodium Chloride Sodium Chloride Sodium Sulphate Sodium Sulphate Magnesium Chloride Sodium Sulphate Manganese Sulphate Sodium Sulphate Magnesium Sulphate Magnesium Sulphate Magnesium Sulphate Calcium Chloride Sodium Sulphate Magnesium Sulphate Calcium Chloride

64 THE CHEMICAL	J FORMULAR1
Mud Bath Salt Ferrous Sulphate 900 g.	Steel (Iron) Baths Formula No. 1
Calcium Sulphate, Precipitated 20 g. Magnesium Sulphate 20 g.	Iron Tartrate 100 g. Distilled Water 900 cc. No. 2
Sodium Sulphate 40 g. Ammonium Sulphate 20 g. Optional, Dry Mud Earth.	Iron Sulphate, Pure 30-60 g. Potassium Carbonate, Pure 120 g. No. 3
"Saltrate Rodell"	Iron Sulphate 30 g. Salt 60 g.
Sodium Chloride, Powder 0.1 g. Magnesium Carbonate 0.5 g. Potassium Carbonate 0.1 g. Lithium Carbonate 0.05 g.	Sodium Bicarbonate 20 g. Sulphur Baths
Calcium Sulphate, Powder 0.25 g. Borax, Powdered 10 g. Sodium Bicarbonate 30.5 g.	Formula No. 1 Potassium Sulphide 50 g. Eau de Cologne 50 g.
Ammonium Carbonate 52.5 g. Sodium Thiosulphate 2.5 g.	Distilled Water 950 ec. No. 2
Sodium Perborate 3 g.	Soft Soap 250 g. Glycerin 50 g. Potassium Sulphide 25 g.
Stimulating Bathing Salt	No. 3
Sodium Chloride, Powder 950 g. Sodium Bicarbonate 50 g. Thyme Oil 2 cc.	Sodium Thiosulphate plus Acid Bath-Water
Bergamot Oil Terpenes 5 cc.	No. 4
Hyme On 2 cc. Bergamot Oil Terpenes 5 cc. Orange Peel Terpenes 1 cc. Bergamot Oil 1 cc. Terpineol 1.5 cc.	a. Sulphur Sublimed 50–100 g. Ammonium Carbonate 950–900 g.
Methyl Naphthyl Ketone 0.5 cc.	b. Distilled Water, Warm 650 cc. Potassium Chromate, Neutral 25-50 g.
Effervescent Tablets for Baths	Mix a , dissolve b , mix both and stir
Formula No. 1 Sodium Bicarbonate 300 g.	several hours, until solid. Press and grind; 120 g. used for a bath.
Sodium Acid Sulphate 275 g. Starch 25 g.	No. 5 (Bain de la Parisienne)
No. 2	Sodium Bicarbonate 870 g.
Saponin, Purified 2 g. Starch 25 g. Sodium Bicarbonate 90 g. Tartaric Acid 70 g. The stability can be increased by	Magnesium Carbonate 10 g. Sulphur Flowers, Ground 100 g. Sulphur, Precipitated 20 g. Selenic Acid 0.1 g.
pressing the bicarbonate and acid separately.	Carbon Dioxide Baths Formula No. 1
Effervescent Tablets with Wetting Agents	Ammonium Carbonate 35 g. Sodium Bicarbonate 20 g. Tartaric Acid 30 g.
(Slow Development of Carbon Dioxide) Formula No. 1	Sodium Perborate 10 g. Sodium Thiosulphate 3 g. Disodium Phosphate 2 g.
Starch 10 g. Sodium Lauryl Sulphonate 10 g. Sodium Bicarbonate 46 g.	No. 2 Sodium Bicarbonate 42 g.
Tartaric Acid 34 g. No. 2	Sodium Acid Sulphate 21 g. Starch 5 g. Sodium Chloride, Powder 30 g.
Sodium Bicarbonate 57 g. Tartaric Acid 38 g.	No. 3
Saponin, Purified 5 g. Stearin, Powder 5 g.	Ammonium Carbonate 25 g. Sodium Bicarbonate 20 g.

	Tartaric Acid	25	
	Sodium Perborate	10	
	Rice Starch	20	
	Manganese Nitrate	1/10	\mathbf{g} .
	Mix all components-except	the p	erbo-
ra	ate—dry and perfume, then	add	the
р	erborate. Press in tablets.		
-			

Mud Bath

Ferrous Sulphate, Crude	900 g
Epsom Salts	20 g
Glauber's Salts	40 g
Ammonium Sulphate	20 g
Gypsum, Crude	20 g
Clay, Dark	50 g
= :	

Foot-Bath Powders (or Tablets) with Perborate

	For	nula
	No. 1	No. 2
Sodium Perborate	170 g.	180 g.
Boric Acid, Powder		60 g.
Borax, Powder	50 g.	
Sodium Acid Car-		
bonate		200 g.
Perfume	5-10 g.	-

Tablet or powder doses for each bath should weigh 10-20 g.

Cold Creams Formula No. 1

T OTHIUM	110. 1	
Cetyl Alcohol		10 g.
Paraffin, Liquid		10 g.
Vaseline, White		80 g.
Water		60 g.

Transparent, soft, white cream.

No. 2

Cetyl Alcohol Paraffin, Liquid Vaseline, White Water No. 3	10 g. 40 g. 50 g. 60 g.
Cetyl Alcohol Paraffin, Liquid Vaseline, White Water No. 4	10 g. 40 g. 15 g. 35 g.
Cetyl Alcohol Paraffin, Liquid Vaseline, White Water	20 g. 20 g. 60 g.

In place of the liquid paraffin there can be used a good vegetable oil. The maximum water-content (37.5%) can be increased by adding 10% wool fat.

Procedure: Melt the fatty materials together and stir, then run in boiling water, a little at a time, not adding ad-

ditional water until previous amount is absorbed.

No. 5

White Beeswax	12 g.
White Petroleum Jelly	12 g.
Peach Kernel Oil	50 g.
Rose Water	25 g.
Borax	1 g.
Perfume	to suit

Grandlagg Cold Cranm

Greaseless Cold Crea	LILL	
Stearic Acid	16	OZ.
Glycerin	48	oz.
Mineral Oil	12	OZ.
Paraffin Wax	2	OZ.
Stronger Ammonia Water	4	OZ.
Water	64	OZ.
Perfume	.75	oz.

Cold Croom

Cold Cream		
1. Diglycol Stearate	14	lb.
2. Paraffin Wax	2	lb.
3. Mineral Oil		gal.
4. Petrolatum (White)		lb.
5. Water	6	gal. fl. oz.
6. Perfume Oil	$5\frac{1}{2}$	fl. oz.
Method of manufacture:		

a. Melt Nos. 1, 2, 3 and 4 at 170° F.
b. Heat 5 to 180° F.

c. Add b to a while mixing. Allow mixer to run until batch is completely emulsified.

d. Allow batch to cool to 125° F. and add 6 and mix at low speed.

e. Batch should be allowed to cool without stirring to 105° F. at which temperature it is poured into jars.

Glycerin Cold Cream

a. Wax, White	80	g.
Spermaceti	80	
Peanut Oil	300	g.
Vaseline	300	g.
Melt.		
b. Glycerin	120	g.
Water	120	g.
Borax	10	
Warm up to 90°; pour into	melte	a.
Add when cool:		
Perfume Composition, Fre	sh	
Odor	20	g.

Triethanolamine Cold Cream (Water-Soluble, Liquid)

a. Paraffin, Liquid Triethanolamine Stearate 14.5 g. Dissolve, warming gently.

b. Water, Distilled	160 g.
c. Perfume When a is dissolved by wa	1.5 g. arming, stir,
and add b slowly. Let star add perfume, then fill into	nd 24 hours.
Cleansing Cream (Semi-Absorbent	
Lanolin	22 g.
White Mineral Oil	25 g.
White Petroleum Jelly Distilled Water	11 g. 42 g.
Perfume	to suit
Cleansing Cream (Non-Absorbent)	
Ceresin	18 g.
White Mineral Oil White Petroleum Jelly	81 g. 1 g.
Perfume	$\begin{array}{ccc} 1 & \mathbf{g.} \\ 0.5 & \mathbf{g.} \end{array}$
Nourishing Crear	n
White Beeswax	9 g.
Spermaceti White Petroleum Jelly	3 g. 35 g.
Benzoated Lard	35 g. 18 g.
Lanolin	4 g.
Liquid Paraffin Distilled Water	9 g. 21 g.
Borax	21 g. 1 g.
Liquid Nourishing C	ream
Lanolin, Anhydrous	16 g.
Stearic Acid Triethanolamine	3 g. 1 g.
Water, Distilled	80 g.
Non-Irritating Crea	ams
U. S. Patent 1,979,	
Formula No. 1	
Vanishing Crean	
Stearic Acid Lanolin (Anhydrous)	220 g. 40 g.
Triethanolamine	40 g. 12.5 g.
Diethylene Glycol Mono-eth	ıyl
Ether Water	75 g. 500 g.
The cream is prepared by	0
acid and lanolin and addin	g them with emaining in-

Water 500 g. The cream is prepared by melting the acid and lanolin and adding them with constant stirring to the remaining ingredients, which are heated to 95° C. An emulsion forms at once which thickens upon cooling. Efficient agitation of the mixture is essential to obtain a smooth product. The solid content, i.e., in No. 1, the lanolin and stearic acid, of a cream of this type may vary from 15% to 35% depending upon the ingredients used and the type of product desired.

No. 2 Cleansing Cream

Stearic Acid 122.5 g.
Lanolin (Anhydrous) 35 g.
White Mineral Oil 210 g.
Triethanolamine 17.5 g.
Diethylene Glycol Mono-ethyl

Ether 40 g. Water 420 g.

The method of preparing this cream is the same as that employed in the previous formula. A cream of this type should have a fairly high content of the ethanolamine in order to completely emulsify the oil so that it may be removed from the skin by washing with water. Various oils and waxes may be used in this type of cream, and the oil content should be fairly high.

No. 3 After Shaving Cream

| Stearic Acid | 15 | g | Triethanolamine | 0.75 | g | Diethylene Glycol Mono-ethyl Ether | 8 | 0.75 | g | Menthol Crystals | 0.75 | g | Ethyl Alcohol (Anhydrous) | 0.5 | g | Water | 75 | g |

The cream is prepared according to the procedure given above. In general, creams of this type are similar to the vanishing creams with the addition of an emollient or a medicant, such as menthol, bay rum, witch hazel or the like.

No. 4

Latherless Shaving Cream

Stearic Acid 350Lanolin (Anhydrous) 67.5 g. White Mineral Oil 169g. Triethanolamine 34 g. Sodium Tetraborate 34 Diethylene Glycol Mono-ethyl 22.5 g. Ether Water 1170

This preparation may be made by the procedure given in No. 1 and the oil may be included in the melted acid and wax mixture which is then added to the other ingredients.

Massage Cream

White Beeswax	12.5	g.
Paraffin Wax	10	g.
White Mineral Oil	50	g.
Distilled Water	26	g.
Borax	1	g.
Perfume	0.5	g.

Massage Preparations

These substances are dispensed in ointment, mixture or solution form, and ap-

plied b	efore vibrat	or tor.	after	trea	ıtment,	usually
		~~	-	- T	-	

Formula No. 1		
Menthol	2.5	g.
Tragacanth	4	g.
Glycerin	12	cc.
	15	cc.
Water 3	00	cc.
No. 2		
Gelatin	2	g.
Water	48	
Glycerin	5	cc
Glycerite of Boroglycerin	45	g.
No. 3		
Fluid Extract of Hamamelis	10	cc.
Wool Fat	60	g.
Petrolatum	30	g.
No. 4		_
Menthol	0.8	g.
Camphor	0.8	g.
		-

Almond Hand-Cleansing Paste

Eucalyptol

Petrolatum

3

96

The "Almond Bran" is made out of two equal parts of sweet and bitter Almonds. One can make a "Glycerin Paste" or a "Camphor Paste."

Glycerin Type

Two hundred fifty pounds of the bran are pounded with 5 lb. of rose water and mixed with the following:

One-quarter pound bean or cornflour, 1-2 chicken eggs, 15 lb. borax, 5 lb. fine potassium carbonate, and about 50 lb. glycerin.

The Camphor Paste is made by adding to the pounded "Almond Bran" a mixture of 25 lb. each of 10% camphor oil and spermaceti, molten together.

After cooling, add a powderized mixture of 100 lb. potato flour and 50 lb. talc, and 100 lb. rose water. Mix well altogether. Color with alkannin or curcuma.

Glycerin Jelly for the Hands

a. { Wheat Starch \ Water \ Glycerin } grind	$10 \\ 15 \\ 100$	g. g. g.
b. Tragacanth, White Alcohol (90%) Methyl-p-Hydroxyben-	2	g.
b. Alcohol (90%)	5	g.
Methyl-p-Hydroxyben-		_
zoate	0.5	g.

Grind a and b separately, mix, warm then on the water bath until odor of alcohol disappears.

Classic II. T. T.		
Glycerin-Honey Jell		
Honey	20	g.
Water	500	g.
Glycerin	450	o.
Amon Amon Cut	15	
Agar-Agar, Cut		4.2
Methyl-p-Hydroxybenzoate	1	g.
Warm to complete swelling	and	solu-
tier manualote if management		
tion percolate, if necessary.	ыш,	and
add:		
Formaldehyde (40%)	1	g.
Perfume Composition	-;	g.
1 ciranic composition	-	8.
Protective Hand Crean	va.ce	
	ns	
Formula No. 1		
Zinc Stearate, U.S.P.	10	O.
Almosterate, O.B.L.	10	g.
Aluminum Subacetate Solu-		
tion N.F. (7½-8%)	15	g.
Gum Camphor Menthol Crystals	3	g.
Menthol Crystals	1	g.
Acid Carbolic, U.S.P.	1/2	g.
Clysonia II C D	1/2	8.
Glycerin, U.S.P. Lanolin, Anhydrous	7/2	
Lanolin, Annydrous	1/2	g.
Gum Tragacanth	$4\frac{1}{2}$	g.
Soap (Low Alkali Content) White Rose Oil Technical	1.8	g.
White Rose Oil Technical	1/2	g.
Triethanolamine	1/2	
	10 72	g.
Water	46	g.
No. 2		
Winn Channels TT CIT	10	
Zinc Stearate, U.S.P.	10	g.
Aluminum Subacetate Solu-		
tion N.F. (7½-8%)	15	\mathbf{g} .
Gum Camphor	3	g.
Menthol Crystals	1	g.
Acid Carbolic, U.S.P.		
Classic II CD	72	g.
Glycerin, U.S.P.	1/2	g.
Lanolin (Anhydrous)	1/2	g.
Gum Tragacanth	41/2	g.
Soap (Low Alkali Content) White Rose Oil Technical	18	g.
White Rose Oil Technical	1/2	g.
Triethanolamine	142	
	4417	g.
Water	441/4"	g.
Sulpho Ammonium	-	
Ichthyolate	2	g.
37 0		_
No. 3		
White Rose Technical Oil	35	er .
		g.
Paraffin Wax	55	g.
Ammonium Sulpho-Ich-		
thyolate	2	g.
Stearic Acid	1	g.
Triethanolamine	1/2	g.
Water	71/2	
	1 72	g.
No. 4		
Glyconyl Monastanata	0	11.
Glyceryl Monostearate		lb.
Magnesium Stearate	14	
Beeswax	3	b.
Petrolatum	10	lb.
Minaral Oil White	E 1	11

Mineral Oil, White

Water

5 lb.

60 lb.

Cuticle Softener Formula No. 1	
White Petrolatum (Short Fiber) Paraffin (mp. 125° F.) Menthol Thymol Color (Oil Soluble Red)	87.75 oz. 9 oz. 3 oz. .25 oz. to suit
Lanolin (Anhydrous) Water (Distilled) Lecithin Cream Petrolatum (Short Fiber) Mineral Oil (White) Perfume	12 oz. 12 oz. 0.5 oz. 55.5 oz. 20 oz. to suit
Skin Cream a. Stearin	85 g.

Lanolin 5 g.
Cetyl Alcohol 10 g.
Melt together.
b. Glycerin (28° Bé.) 36 g.
Triethanolamine 5 cc.
Borax knifepointful
Water 250 cc.
Boil.

Add b slowly to a, stir until cold. Perfume as desired is added at the end.

"Penetran" Skin Cosmetic

20	cc.
25	cc.
	g.
2.5	g.
47	cc.
	25 5 0.5 2.5

Wrinkle "Removing" Creams

Lanolin anhydrous 20 (parts by weight), cocoa butter 10, stearin 10, olive oil 12, cholesterol 2, lecithin 4, water 60, moldex 0.4, sodium benzoate 1. According to another method, a melted base is first prepared with white wax 60 (grams), spermaceti 10, stearin 50, lanolin 60, cocoa butter 40, and sweet almond oil 180. In this melt are dissolved 1.2 grams cholesterol, with further addition, after complete solution, of 170 g. water, 1.5 g. sodium benzoate and moldex, the mass being stirred until it thickens.

Skin "Food" Formula No. 1

Lanolin (Anhydrous)		
U.S.P.	36.4	g.
Spermaceti, U.S.P.	6.4	g.
Snow White Petrolatum	1,	-
U.S.P.	48.2	g.

Distilled Water Perfume Oil	7.875 g. 1.125 g.
No. 2	
Almond Oil	24 g.
Lanolin	22 g.
Soft Paraffin	11 g.
White Beeswax	3 g.
Rose Water	40 g.
Perfume	to suit

Mosquito Repelling Cream Formula No. 1

a. Wheat Starch	5 g.
". (Water	10 g.
b. Glycerin (28° Bé.)	45 g.
c. Lanolin	30 g.
d. Clove Oil	5-10 g.

Grind a until homogeneous, add b, and warm gently until a homogeneous jelly is formed. Cool, and grind now with c and d in a mortar very thoroughly until distribution is satisfactory. Fill at once into collapsible tubes.

No. 2

(White Wax	50	g.
a. {White Wax Spermaceti	5 0	g.
	4	g.
b. Borax (0.96)	40	g.
c. Water Wheat Starch Gelatin Sodium Benzoate	510	cc.
Wheat Starch	1	g.
c. Gelatin	4	g. g.
Sodium Benzoate	0.5	g.

Make up cream as usual pouring b into a, then add the solution c which is to be made up before (soak cold, then warm to clear solution, if necessary, pour through a fine sieve), stir thoroughly, stop heating, stir until cooled, and add

Eucalyptus Oil 50 cc.

No. 3

Eucalyptus Oil		0.5	cc.
Caryophyllum Oil		0.5	
Lavender Oil		0.5	cc.
Quinine Sulphate		1	g.
Glycerin Salve	to make	100	g.

No. 4

Soap Solution 2.5-2	g.
Glycerin 4 To this cream add:	ΰg.
	ŏğ.
M4h1 1	
MEHINI	g.
Sodium Benzoate 1	g.
Citronella Oil 1	cc.
Caryophyllum Oil 0.5	cc.
Alcohol 10	cc.
Tincture of Green Soap 10	cc,

	DIRITED IDIOON
Mosquito Repellants	Alcohol 12 g.
Formula No. 1	Berswax, White 8 g.
Pyrethrum Flowers 10 g.	Lanolin (Anhydrous) 8 g.
Isopropyl Alcohol, or Ethanol	Glycerin 60 g.
with Thymol 100 g.	Water 830 g.
Oil of Cloves 2 g.	Beta Naphthol 1 g.
No. 2	Essential Oils as in Formula No. 1
Oil of Eucalyptus 45 g.	Treatment as in No. 1, saponify the
Oil of Thuia 20 g.	fats (wax, lanolin, stearin) together.
Oil of Laurel $5 g$.	No. 3
Phenol 3 g.	a. Agar-Agar 2.5 g.
Camphor 20 g.	Glycerin 100 g.
Alcohol 100 g.	Water 750 g.
Turpentine Oil 50 g.	b. Glyceryl Monostearate 120 g. Spermaceti 100 g.
Quassia, Tincture 40 g.	Spermaceti 100 g.
Pyrethrum Extract 50 g.	Melt.
Xylol to make 1000 cc.	Pour a hot into b , make emulsion, stir.
No. 3	Add boiling water up to 980 g. Add,
Pyrethrum Extract 0.5 g.	when cold:
Amyl Salicylate 3.5 g.	Moldex or Other Good
Petroleum (bp. 182-292°;	Preservative 2 g.
sp. gr. 0.801) 96 g.	Essential Oils 12 g.
No. 4	(See Formula No. 1)
Pyrethrum Powder 1 g.	
Derris-Root Powder 1 g.	All Weather Cream
Tobacco Powder 0.5 g.	
Derris-Root Powder 1 g. Tobacco Powder 0.5 g. Alcohol, Diluted 25 g.	a. Stearic Acid 210 g. Adeps Lanae, Anhydrous 50 g.
Percolate thoroughly and filter; add:	
oil of eucalyptus or menthol to suit.	$ \begin{cases} \text{Glycerin} & \text{133 g.} \\ \text{Triethanolamine} & \text{20 g.} \end{cases} $
	b. Triethanolamine 20 g. Borax 5 g.
Mosquito Protection Cream	Distilled Water 582 cc.
(Non-Greasy)	Melt up a to about 65° C., add b boil-
Formula No. 1	ing hot, in thin jet, stirring thoroughly
Soak	until cold.
a. Agar-Agar 2 g.	Printer of the state of the sta
Water, Cold 400 g.	Night Cream (Greasy)
Then warm slowly over gentle heat:	(Paratin Oil White 9500 -
b. Melt Stearin 60 g.	Paraffin Oil, White 2500 g.
c. Alcohol (95%) 10 g.	a. Beeswax, Bleached 500 g.
d Potassium Carponate 6 g.	a. Wax, Scale 500 g. Beeswax, Bleached 500 g. Adeps Lanae, Anhydrous 500 g.
(Water 440 g.	Distilled Water 3000 cc.
Glycerin (28° Bé.) 68 g.	b. Triethanolamine 75 g.
Make up emulsion by warming and	b. Triethanolamine 75 g. Borax 35 g.
stirring.	Melt a together at 75° C.; add b which
Add a to the emulsion of $b - c$ in d ,	is at same temperature, to a. Stir until
both should be 80° C.; stir continously. When cold, add 12 g. of the following	cold.
mixture:	Annual an
	Non-Greasy Cream
Citmonalla Oil	Formula No. 1
Comphen	
Eucalyptus Oil 4.5 g.	Stearie Acid 230 g.
Alcohol 7 g.	a. Wax, Scale 40 g. Adeps Lanae, Anhydrous 10 g.
No. 2	Clysopin
Treatment as above:	
Agar-Agar 2.2 g.	
Stearin 60 g.	Borax 5 g. Distilled Water 562 cc.
Potassium Carbonate 4 g.	Melt a and warm up b in another con-
Sal Soda 2 g.	tainer. Mix both (a and b should be 65°
	- Co and o should be 65

C. boiling) pouring b into a in thin jet. Stir until cold.

No. 2

	Stearic Acid Adeps Lanae, Anhydrous	170 g.
	Adeps Lanae, Anhydrous	13 g.
a.	Wax, Scale	13 g.
	Spermaceti	5 g.
	Wax, Scale Spermaceti Cetyl Alcohol	4 g.
	Glycerin (28° Bé.) Triethanolamine	80 g.
	Triethanolamine	13 g.
b	Borax	5 g.
	Borax Distilled Water	697 ec.

Melt up waxes $(65-70^{\circ})$, add b hot (boils) in thin jet, stirring thoroughly. Optionally, 100 water may be substituted by witch hazel (1:1). Stir until cold.

Liquid Cream

Stearic Acid	50	g.
Adeps Lanae, Anhydrous		ğ.
a. Cetyl Alcohol	1	g.
a. Adeps Lanae, Anhydrous Cetyl Alcohol Beeswax	1	g.
	20	g.
Glycerin Triethanolamine	2	ğ.
b. Borax	2	g.
Witch Hazel (1:1) Distilled Water	- 75	g.
Distilled Water	625	cc.

Melt up together a at 60-70° C. Heat b to boiling, then add in thin jet, stirring vigorously, to a. Stir until cold.

To all above-mentioned creams, perfume should be added during cooling (0.5-0.7%). The perfume components should be colorless, and should not irritate the skin. No alcoholic compositions should be used.

Turtle Oil Cream

1.	Diglycol Stearate	14	lb.
2.	Mineral Oil	33/4	gal.
	Lanolin	6	lb.
4.	Petrolatum (White)	2	
5.	Water	6	gal. fl. oz.
6.	Turtle Oil	$5\frac{1}{2}$	fl. oz.
7.	Perfume Oil	51/2	fl. oz.
8.	Solution Yellow Color		
	Made by Dissolving		
	Yellow Dye 2 drams in		
	Mineral Oil 14 fl. oz.	81/4	fl. oz.

Method of manufacture:

a. Melt 1, 2, 3, 4, 6 and 8 at 170° F.
b. Heat 5 to 180° F.

c. Add b to a while mixing. Allow mixer to run until batch is completely emulsified.

d. Allow batch to cool to 125° F. and add 7, and mix at low speed.

e. Batch should be allowed to cool without stirring to 100° F. at which temperature it is poured.

Boro-Glycerin Lanolin Cream

$a. egin{cases} ext{Boric Acid} \ ext{Glycerin} \ ext{Water} \end{cases}$	10	g.
a. Glycerin	40	g.
Water	250	g.
Dissolve.		
b. Lanolin, Anhydrous Vaseline, White	100	g.
Vaseline, White	600	g.
Melt gently.		
c. Rose Oil, Artificial	10	cc.
or Eau de Cologne Oil	20	ec.

Tragacanth-Glycerin Base (Used Below)

Tragacanth, White, Fine

Powder Glycerin 5 g. Grind thoroughly in mortar and add:

Water, Warm Add while stirring and in small portions, warm up to 40° C. Stir until paste is homogeneous.

Menthol Cream

Menthol Moldex or Other Good Pre-	0.2	g.
servative Perfume Oil	0.2	g.
Alcohol (95%)	0.3 5	
Dissolve and add Glycerin	5	g.
Add above made Tragacanth-Glycerin	100	g.

Lemon Juice Cream U. S. Patent 1,990,676

Five parts of oxy-cholesterin and 95 parts of petrolatum are thoroughly mixed to form an absorption base. Twenty parts of petrolatum and three parts of beeswax are melted together, and 30 parts of the base are added with thorough stirring. Fifty parts of nat-ural lemon juice are added to the above mixture while still hot and stirring is continued until the mass is cool.

Ink Removing Cream U. S. Patent 1,968,304

A substantially non-aqueous cream for the removal of ink stains from the skin contains about 500 g. of zinc stearate, about 300 g. of citric acid, about 500 cc. of 95 per cent ethyl alcohol and about 2000 cc. of diethylene glycol.

Deodorant Cream Formula No. 1

Benzoic Acid	4 g.
Zinc Oxide	12 g.
Lanolin	4 g.
Petrolatum (Snow White)	80 g.
Perfume	to suit
N_0 2	

British Patent 425,059

Coconut Oil	63 g.
Lemon Oil	5.2 g.
Boric Acid, Powdered	21 g.
Starch, Powdered	10.5 g.
Lanolin	0.2 g.
Perfume	0.1 g.
No. 3	
77	1 00

Formaldehyde 1 oz. Vanishing Cream 99 oz.

Powder Cream Base

Using specified quantities, preparation of the cream base may proceed on the following lines: A mixture of about 500 g. distilled water, 20 g. potassium carbonate and 125 g. glycerin is heated almost to boiling point in a capacious vessel constructed of well enamelled mate-Two hundred grams stearic acid melted in another vessel are cautiously introduced, a little at a time, into the hot potassium carbonate solution. Violent carbon dioxide evolution ensues and continues until the last portion of stearic acid has been added. When gas development ceases, indicating completion of the reaction, heating is discontinued and the batch transferred to another vessel fitted with stirring gear. An additional 1000 g. water and 125 g. glycerin are added and the mix stirred until cold and viscous. Cold-stirring is important for securing a fine, uniform emulsion and for preventing settlement of stearic acid particles. Certain variations in preparation can be practiced, such as replacement of glycerin by white liquid paraffin or addition of 125 g. groundnut oil to facilitate emulsification.

Ruggles' Cream

00	
Powdered Stearic Acid	75 g.
Potassium Carbonate	15 g.
Distilled Water	320 g.
Powdered Borax	5 g.
Quince Jelly	75 g.
Distilled Water	100 g.
Powdered Zinc Oxide	10 g.
Glycerite Starch	400 g.

Melt the stearic acid. At the same time dissolve the potassium carbonate in 320 cc. of distilled water and heat to about 170° F. on water bath. Bring stearic acid to the same temperature and mix them. Continue this temperature on the water bath, with occasional stirring, until the reaction is perfectly complete.

Dissolve the powdered bornx in 100 cc. of distilled water, add the quince jelly and heat on water bath to about 170° F. Add this mixture to the first, which should be at the same temperature, and again leave on water bath until reaction is complete.

Heat the glycerite of starch to the same temperature, stir in the powdered zine oxide with a glass stirring rod and add to the other mixture, stirring occasionally.

Let cool and add perfume (oil ylang

ylang recommended).

The most important essential is to employ a perfect glycerife of starch. Use Kingsford's or other suitable grade of corn starch and U. S. P. Glycerin and make it up fresh for each batch.

It is also essential to have all three batches at exactly the same temperature

when mixing them.

Skin Oil with Isocholesterin Paraffin Oil plus Preserved Fatty Oil 97 cc. Isocholesterin, Technically Pure 3 g. or Same, Chemically Pure 2 g.

Skin Oil with Lanolin Lanolin, Bleached 5 g. Paraffin Oil or Fatty Oils 95 cc.

Skin Oil with Wool Wax
Wool Wax, Bleached, Purified 5 g.
Fatty Oil 35 cc.
Paraffin Oil 60 cc.

Skin Oil with Cetyl Alcohol
Cetyl Alcohol, Pure 3-5 g.
Paraffin Oil plus Fatty Oil,
Preserved (1:1) 97-95 cc.

Skin Oil with Triethanolamine Oleate Triethanolamine Oleate, Pure 2 g. Fatty Oil 98 cc.

Non-Irritating Skin Oil
Diglycol Laurate Neutral 4 g.
Olive Oil 96 cc.
Perfume to suit

72 THE CHEMICA	L FORMULARY
Lecithin Skin Oil Formula No. 1 Lecithin from Eggs 10-30 g.	Witch Hazel Skin Oil Witch Hazel Leaves, Powder 100 g. Fatty Oil, Preserved 900 cc.
Paraffin Oil 170–190 cc. Olive Oil, Preserved 800 cc. Perfume, to suit 5 g.	Pour hot oil over leaves, let stand for 8 days. Filter.
No. 2	Massage Oil
Lecithin from Brain Sub- stance 20 g. Paraffin Oil 180 cc. Olive or Peanut Oil, Pre- served 800 cc.	Paraffin Oil 75 cc. Parachol (Absorption Base) 5 g. Olive Oil, Preserved 20 cc.
	Muscle Oil
Skin Oil "Huile Ambrosiaque" Ambergris, Best Quality 10 g. Behen Oil 990 cc.	Castor Oil, Deodorized 66.6 cc. Alcohol (92–95%) 33.3 cc. Cholesterin, Pure 0.1 g.
Perfume to suit Grind the amber with glass-powder and introduce into the warmed oil. Shake well. Filter after 3-4 weeks.	Sport Oil (for Swimmers) Octadecyl Alcohol (Pure) 5 g. Fatty Oil, Preserved 55 cc. Paraffin Oil 40 cc.
Skin Oil with Wool Fat Alcohols	Chalastavia Oil
Parachol	Cholesterin Oil
(Absorption Base) 5–10 g. Paraffin Oil 95–90 cc.	Fatty Oil, Pure, or in Mixture with Paraffin Oil 1000 cc. Cholesterin, C.P. 5-10 g.
Skin Cleansing Oil Parachol or Absorption Base 2 g. Triethanolamine Oleate, Pure 0.5 g. Fatty Oil, Preserved 97.5 cc. Add a little Triethanolamine.	Cholesterin-Lecithin Oil Same as Cholesterin Oil, but besides add Lecithin (Eggs, Brain-Substance) 20-30 g.
	Face Lotions
Skin Nourishing Oil	Formula No. 1
Egg Oil 5 g. Parachol (Absorption Base) 5 g. Lecithin 1 g. Sperm (Whale) Oil, Genuine, Deodorized 20 cc. Fatty Oil, Preserved 69 cc.	
Skin "Stimulating" Oils	Triethanolamine 0.5 cc. Glycerin 4 cc.
Formula No. 1	Alcohol (30%) 95.5 cc.
Parachol (Absorption Base) 5 g. Oxycholesterin, Artificial 3 g.	Perfume to suit
Fatty Oil (Olive, Sesame, Peanut), Preserved 92 cc.	Orange Flower Water 800 cc.
	Eau de Cologne 200 cc.
No. 2	Triethanolamine 6 cc. Spirits of Camphor 20 cc.
Parachol (Absorption Base) 5 g.	Glycerin 100 cc.
Cetyl Alcohol, Pure 3 g. Fatty Oil, Preserved 91 cc.	No. 4
2 333, 2 20001100	Camphor 20 cc. Alcohol (96%) 850 cc.
	Alcohol (96%) 850 cc. Glycerin (28° Bé.) 50 cc.
Astringent Skin Oil	Perfume 30 cc.
Aluminum Stearate 3 g. Fatty Oil 97 cc.	Distilled Water 1500 cc. Triethanolamine 15 cc.

No. 5		Alcohol	6 g.
Triethanolamine	5 cc.	Glycerin, C.P.	3 g.
Alcohol (96%)	500 cc.	Almond Oil	10 g.
Spirits of Camphor	100 cc.	Distilled Water about	t 85 g.
Perfume	10 cc.		
Glycerin	20 cc.	Face Lotion (For Oily	Skin)
Witch Hazel, Distilled	1000 cc.	Sulphur, Precipitated	2 g.
		Classin CD	5 g.
For Dry Skin: No	0. 6	Glycerin, C.P.	3 g.
Mineral Oil, White	35 cc.	Camphor Spirit (10%)	10 g.
Beeswax	20 g.	Lavender Water	1 g.
Amino Stearin	8 g.	Borax	21 6
Water	50 cc.	Distilled Water	81 g.
Warm together and mi	x vigorously		
until emulsified.		Acne Face Lotion	ì
No. 7		Formula No. 1	
Vaseline Oil	72 cc.		
Amino Stearin	14 g.	Acetic Acid (96%) or	5 g.
Water	200 cc.	Benzoic Acid	500 8.
No. 8		Alcohol (95%)	500 g.
	5 cc.	Lavender Oil	4 g.
Triethanolamine	0.0	Water	466 g.
Aromatic Spirit	30 cc. 12.5 cc.	Glycerin (28° Bé.)	25 g.
Bergamot Oil	0.5 cc.	Let stand several weeks. I	filter.
Oil Orange Flowers	2 cc.	No. 2	
Lemon Oil	15 cc.	Potassium Soap from	
Rosemary Oil		Olive Oil (Neutralized)	100 g.
Alcohol (70%)	940 cc.	Alcohol (90%)	500 g.
No. 9		Lavender Oil	5 g.
Camphor	25 g.	Rose Oil, Artificial	5 g.
Alcohol	850 cc.	Water	390 g.
Glycerin	25 cc.	11 2001	Sh., 8.
Perfume Mixture	30 cc.	77 777. 4	
Distilled Water	1570 cc.	Face Water	
No. 10		Triethanolamine	0.5 g.
Borie Acid	10 g.	Glycerin	4 g.
Glycerin	29 cc.	Alcohol	33 g.
Menthol	1 g.	Perfume	0.5 g.
Perfume	5 cc.	Distilled Water	62 g.
Alcohol	255 cc.		
Hamamelis Distillate	300 cc.	Prophylactic Face Wa	itora
Rose Water	400 ec.		
No. 11		Formula No. 1	
Alcohol	450 cc.	Ammonium Chloride, C.P.	0.5 g.
Camphor, Spirits of	100 cc.	Witch Hazel	20 ee.
Perfume	10 ec.	Rose Water	10 ec.
Hamamelis Distillate	440 cc.	Distilled Water	69.5 cc.
No. 12		No. 2	
	100	Ammonium Chloride	2.5 g.
Potassium Carbonate	400 g.	Cherry Laurel Water	10 ce.
Distilled Water	2000 cc.	Witch Hazel	10 cc.
Orange Flower Water	1000 cc.	Rose Water	20 ec.
Alcohol	100 cc.	Distilled Water	57 ec.
Perfume	to suit	Diethylene Glycol	0.5 cc.
No. 13	50	- 10011/1020	
Borax	50 g.	Name and the second sec	
Sodium Thiosulphate	500 g.	Kummerfeld's (Face)	Water
Distilled Water	8500 cc.	Sulphur, Colloidal, or Fine	ly
Glycerin	500 cc.	Precipitated	2 g.
Eau de Cologne	500 cc.	Glycerin	12 cc.
		Spirits of Camphor	4 cc.
Face Lotion (For Dry	Skin)	Eau de Cologne	20 cc.
Lanolin or Cholesterol	0.05 g.	Distilled Water	100 cc.
Lecithin	0.05 g.	Optionally: Addition of Bo	
ALLEU TOTAL	5. 5. E.	- paronomy . reddiction of 190	111 1 111

Sulphur Face Water Sulphur, Colloidal 3 g. Potassium Carbonate 1.5 g. Glycerin 5 cc. Spirits of Camphor 4 cc. Alcohol 10 cc. Distilled Water 76.5 cc. Skin Lotion Gum Tragacanth 4 oz. Glycerin 3 oz. Phenol 1 oz. Oil of Teel 120 oz. Water 360 oz. Perfume 2 oz. Modern Glycerin-Sulphur Lotion Colloidal Sulphur in Glycerin (24%) Tincture of Green Soap 100 g. Eau de Cologne—Oil 1 g. Water, Distilled 799 g. Glycerin and Cucumber Lotion Cucumber Perfume 5 g. Alcohol (95%) 50 g. b. Senzoic Acid 0.3 g. Cucumber Perfume 5 g. Glycerin and Cucumber Lotion Cucumber Perfume 5 g. Tragacanth, Fine, White 5 g. Glycerin c together, then add a and b in small portions, grinding to get homogene-	oz. gal. gal. ay be
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Spirits of Camphor 4 cc. Alcohol 10 cc. Distilled Water 76.5 cc. Distilled 76.	oz. gal. gal. ay be
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Oil of Teel 120 oz. Water 360 oz. Perfume 2 oz. Witch Hazel 550 oz. Perfume 2 oz. Modern Glycerin-Sulphur Lotion Colloidal Sulphur in Glycerin (24%) 100 g. Tincture of Green Soap 100 g. Eau de Cologne—Oil 1 g. Water, Distilled 799 g. Glycerin and Cucumber Lotion Cucumber Perfume 5 g. Alcohol (95%) 50 g. Benzoic Acid 0.3 g. Cucumber Perfume 5 g. Glycerin 100 g. Grind c together, then add a and b in small portions, grinding to get homogene-	
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Perfume 2 oz. Modern Glycerin-Sulphur Lotion Colloidal Sulphur in Glycerin (24%) 100 g. Tincture of Green Soap 100 g. Eau de Cologne—Oil 1 g. Water, Distilled 799 g. Glycerin and Cucumber Lotion Cucumber Perfume 5 g. Alcohol (95%) 50 g. b. $\begin{cases} Alcohol (95\%) & 50 g. \\ Benzoic Acid & 0.3 g. \\ Cucumber Perfume & 5 g. \\ Glycerin & 100 g. \\ Glycerin & 100 g. \\ Grind c together, then add a and b in small portions, grinding to get homogene- Modern Neutral Face Water Alcohol (40%) 920 Diethylene Glycol 30 Glycerin, C.P. 50 Alcohol (40%) 920 Diethylene Glycol 30 Glycerin, C.P. 50 Strick Alcohol (40%) 920 Diethylene Glycol 30 Glycerin, C.P. 50 Strick Alcohol (40%) 920 Clycerin, C.P. 50 Strick Alcohol (40%) 920 Clycerin,$	g.
Modern Glycerin-Sulphur Lotion Colloidal Sulphur in Glycerin (24%) Tincture of Green Soap Eau de Cologne—Oil Glycerin and Cucumber Lotion Cucumber Perfume 5 Benzoic Acid Cucumber Perfume 5 Cucumber Perfume 5 Benzoic Acid Cucumber Perfume 5 Cucumber Perfume 5 Benzoic Acid Cucumber Perfume 5 Glycerin Cucumber Perfume 6 Gl	cc.
Modern Glycerin-Sulphur Lotion Colloidal Sulphur in Glycerin (24%) Tincture of Green Soap Eau de Cologne—Oil Glycerin and Cucumber Lotion Cucumber Perfume 5 Benzoic Acid Cucumber Perfume 5 Cucumber Perfume 5 Benzoic Acid Cucumber Perfume 5 Cucumber Perfume 5 Benzoic Acid Cucumber Perfume 5 Glycerin Cucumber Perfume 6 Gl	g.
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Constitute Surplier in Glycerin 100 g. Tincture of Green Soap 100 g. Eau de Cologne—Oil 1 g. Water, Distilled 799 g. Glycerin and Cucumber Lotion Cucumber Perfume 5 g. $Alcohol (95\%)$ 50 g. $b.$ $Benzoic Acid 0.3 g. Cucumber Perfume 5 g. c. Tragacanth, Fine, White 5 g. c. Tragacanth, Fine, White 5 g. c. Glycerin 100 g. c. Grind c. together, then add a and b in small portions, grinding to get homogene-$	
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Eau de Cologne—Oll 1 g. Water, Distilled 799 g. Water, Distilled 799 g. Glycerin and Cucumber Lotion Cucumber Perfume 5 g. $\{ Alcohol (95\%) \\ Benzoic Acid 0.3 g. \\ Cucumber Perfume 5 g. \\ C. Tragacanth, Fine, White 5 g. Glycerin 100 g. Grind c together, then add a and b in small portions, grinding to get homogene- Face Water for Mottled Skin or F Zinc Sulphate, C.P. 1 Citric Acid, C.P. 0.4 Hydrogen Peroxide (3-10\%) 89.5 Freckle Lotion Dissolve: Potassium Carbonate 66 Potassium Chlorate 22 Sugar 66$	g.
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Glycerin and Cucumber Lotion Cucumber Perfume 5 g. Alcohol (95%) 50 g. Benzoic Acid 0.3 g. Cucumber Perfume 5 g. Cucumber Perfume 5 g. Tragacanth, Fine, White 5 g. Glycerin 100 g. Grind c together, then add a and b in small portions, grinding to get homogene- Zinc Sulphate, C.P. 1 Citric Acid, C.P. 0.3 Hydrogen Peroxide (3-10%) Freckle Lotion Dissolve: Potassium Carbonate Potassium Chlorate 20 Borax 13 Sugar 66	eckles
$b. \begin{cases} \text{Aconor} & (95\%) & 50 & \text{g}. \\ \text{Benzoic Acid} & 0.3 & \text{g}. \\ \text{Cucumber Perfume} & 5 & \text{g}. \\ \text{Cucumber Perfume} & 5 & \text{g}. \\ \text{Glycerin} & 100 & \text{g}. \\ \text{Glycerin} & 100 & \text{g}. \\ \text{Grind c together, then add a and b in small portions, grinding to get homogene-} \end{cases} $ $Borax \qquad 14$ $Sugar \qquad 60$	or
$b. \begin{cases} \text{Aconor} & (95\%) & 50 & \text{g}. \\ \text{Benzoic Acid} & 0.3 & \text{g}. \\ \text{Cucumber Perfume} & 5 & \text{g}. \\ \text{Cucumber Perfume} & 5 & \text{g}. \\ \text{Glycerin} & 100 & \text{g}. \\ \text{Glycerin} & 100 & \text{g}. \\ \text{Grind c together, then add a and b in small portions, grinding to get homogene-} \end{cases} $ $Borax \qquad 14$ $Sugar \qquad 60$	g.
$b. \begin{cases} \text{Aconor} & (95\%) & 50 & \text{g}. \\ \text{Benzoic Acid} & 0.3 & \text{g}. \\ \text{Cucumber Perfume} & 5 & \text{g}. \\ \text{Cucumber Perfume} & 5 & \text{g}. \\ \text{Glycerin} & 100 & \text{g}. \\ \text{Glycerin} & 100 & \text{g}. \\ \text{Grind c together, then add a and b in small portions, grinding to get homogene-} \end{cases} $ $Borax \qquad 14$ $Sugar \qquad 60$	۶.
$b. \begin{cases} \text{Aconor} & (95\%) & 50 & \text{g}. \\ \text{Benzoic Acid} & 0.3 & \text{g}. \\ \text{Cucumber Perfume} & 5 & \text{g}. \\ \text{Cucumber Perfume} & 5 & \text{g}. \\ \text{Glycerin} & 100 & \text{g}. \\ \text{Glycerin} & 100 & \text{g}. \\ \text{Grind c together, then add a and b in small portions, grinding to get homogene-} \end{cases} $ $Borax \qquad 14$ $Sugar \qquad 60$	ec.
c. Tragacanth, Fine, White 5 g. Glycerin 100 g. Potassium Carbonate 20 Grind c together, then add a and b in small portions, grinding to get homogene- Sugar 66	
c. Tragacanth, Fine, White 5 g. Glycerin 100 g. Potassium Carbonate 9 Potassium Chlorate 20 Potassium Chlorate 11 Borax 11 Small portions, grinding to get homogene-	
Glycerin 100 g. Potassium Chlorate 20 Grind c together, then add a and b in small portions, grinding to get homogene- Sugar 60	
Grind c together, then add a and b in small portions, grinding to get homogene- Sugar 66	g.
small portions, grinding to get homogene- Sugar Sugar 60	g.
sman portions, grinding to get nomogene-	g.
	g.
The state of the s	
Cucumber and Egg Lotion Rose Water 336 Orange Flower Water 356	g. ec.
Cucumber and Egg Lotion Orange Flower Water 355 Cucumber Juice 400 c Glycerin 150	cc.
Alcohol 50 g. differni	00.
Cucumber Juice 400 g. Glycerin 150 Alcohol 50 g. Skin Cleansing Lotion Benzoic Acid 0.25 g. Skin Cleansing Lotion Egg Yellow 1-2 g. British Patent 423,426 Lavender Oil 3 g. Sodium Biborate 1.3 Rose Oil, Artificial 1 g. Sodium Biborate 1.3 Glycerin 100 g. Potassium Alum 2.3	
Egg Yellow 1-2 g. Details Details 192 400	
Lavender Oil 3 g. British Patent 423,426	
Rose Oil, Artificial 1 g. Sodium Biborate 1.3	g.
Glycerin 100 g. Potassium Alum 2.3	g.
Rose Oil, Artificial 1 g. Sodium Biborate 1.3 Glycerin 100 g. Potassium Alum 2.3 Soda Ash 1.7 Water 100	g.
Alcohol (45%) 900 cc. Evaporate down to half of volu	ne.
Tri- (or Di-) Ethylene Glycol 30 g	
Citric Acid 5 g. Liquid Deep Pore Cleanser	
orycerin 50 g. Witch Hazer Extract, C.S.F. 50	
witch Hazel 35 cc. Alcohol 28	oz.
Polyalkyl-glycol Ether	
Face Water, Astringent (Glycopon S)	oz.
Alcohol (35%) 950 cc.	oz.
Diethylene Glycol 30 g. Face Pack	oz.
Glycerin 15 g. Put on face for 20 min. a mixt	OZ.
Tannic Acid, Pure 3 g. Oat Flour	oz. oz.
Phosphoric Acid, C.P. 2 g. Arnica Flowers	oz. oz.

		1
Chamamila Flawara	2 g	.
Chamomile Flowers		
Hamamelis Leaves		
Rosemary Leaves Camphor Water		
Camphor Water	30 ce	-
Treat afterwards with as	tringent	lo-
tion of	0.05	
Tannic Acid	0.25 g	
Rose Water	25 g	
Hamamelis Water	50 g	
Orange Flower Water	25 g	. 1
		1
Hand Lotion		
Formula No. 1		
Alcohol, Ethyl	600 ce	.
	100 cc	
Glycerol		
Menthol	5 g.	
Perfume, Rose Oil, Etc. Salicylic Acid	1 cc	
Salicylic Acid	2 g.	
Water	300 cc	
No. 2		1
Alcohol, Ethyl	550 cc	. (
(flycerol	175 ee	. 1
Menthol	3 g.	
Dorfung or desired about	1 cc	
Perfume, as desired, about Salicylic Acid	2 g	- 1
	2 g.	. 1
Water	275 ce	
No. 3		1
Alcohol, Ethyl	500 cc	
Glycerol	250 ee	
Menthol	1 g.	. 1
Perfume, as desired, about Salicylic Acid	1 ee	.
Salievlie Acid	2 g.	
Water	250 cc	. 1
A lavender coloration of	varying	ın-
tensity may be obtained by a	dding tra	ces
of ferric chloride solution.	r'ormula .	No.
3 gives a rather oily lotion.		
		- 1
Tow Cost Almond T.	.tian	-
Low Cost Almond Lo		1
1. Diglycol Stearate	7 lb.	l
	0 gal.	1
3. Gum Tragacanth So-		Į
lution	6 gal.	-
4. Benzaldehyde	3 fl. oz	.
5. Oil of Bergamot	11/2 fl. oz	
	- /2	1
Method of manufacture:		
a. Melt No. 1 at 160° F.		1
b. Heat No. 2 to 205° F. a	ınd run i	nto
stone jar (note final ten	aperature	of
water after dumping in	to jar m	ust
not be below 170° F)	J 1	
stone jar (note final ten water after dumping in not be below 170° F.). c. With high speed agitat	ar manni	nor
odd a (molton of 1000	muni runin	g,
add a (molten at 160° at least 170° F. and all	r.) to b ,	at
at least 170° F. and all	ow mixer	to
run until temperature ha	s dropped	to
140° E		1

d. Add 3 to batch while mixture is still

c. Add 4 and 5 immediately after 3 and allow mixer to continue running

140° F.

running.

until temperature has dropped to 90° or 95° F.

The gum solution is made as follows: Gum Tragacanth $2\frac{1}{2}$ lb. Water 50 gal.

Allow the gum to soak for several hours and beat into solution.

Rose Lotion

 1. Diglycol Stearate
 7
 lb.

 2. Water
 30
 gal.

 3. Gum Solution
 6
 gal.

 4. Oil of Rose
 3
 fl. oz.

5. Red Color Solution
Made by Dissolving
Red Dye, 1 oz., in
Water, 1 qt. 34 fl. oz.

Method of manufacture:

a. Melt No. 1 at 160° F.
b. Heat No. 2 to 200° F. and run into stone jar (note: final temperature of

water after dumping into jar must not be below 170° F.).

c. With high speed agitator running add a (molten at 160° F.) to b at at least 170° F. and allow mixer to run until temperature has dropped to 140° F.

d. Add 3 to batch while mixer is still

running.

e. Add 4 and 5 immediately after 3 and allow mixer to continue running until temperature has dropped to 90° or 95° F.

The gum solution is made as explained under almond lotion.

Lemon Lotion

Diglycol Stearate
 Water
 Box 10.
 July 2.
 July 30.
 July

3. Gum Solution 6 gal. 4. Oil of Lemon 1½ fl. oz.

5. Yellow Dye 34 oz.

Method of manufacture:

a. Melt No. 1 at 160° F.

b. Heat No. 2 to 200° F, and run into stone jar (note: final temperature of water after dumping into jar must not be below 170° F.).

c. With high speed agitator running add a (molten at 160° F.) to b at at least 180° F. and allow mixer to run until temperature has dropped

to 145° F.

d. Add 3 to batch while mixer is still

running.

e. Add 4 and 5 immediately after 3 and allow mixer to continue running until temperature has dropped to 95° or 100° F.

The gum solution is made as explained under almond lotion.

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Milky Lotion with Pec	tin	Borax	20	g.
Base Emulsion (See Below)		c. dissolved in	000	
Distilled Water	445 g.	Water, Warm	880	g.
Perfume Base Emulsion	5 g.	Add c cold to a and b.		
	710 m	D / 01 17 17	•	
Distilled Water Mineral Oil	710 g. 180 g.	Dusty Odor Face Lot	nons	
Dried Pectin	50 g.	Formula No. 1	1	00
Citric Acid	10 g.	Glycerin Lactic Acid		cc.
Extract Chamomile Flowers	50 g.	Menthol	0.5	
Moisten the pectin with a li		Opoponax—Perfumes with		
nd then rub with a little water		Violet Root Oil, etc.	0.5	
he citric acid is dissolved u nucilage is obtained. The pe	etin swells	Alcohol (35%)	$0.3 \\ 97.5$	
o a large extent. In the r	est of the	No. 2	01.0	00.
rater dissolve the liquid char	momile ex-	Glycerin	1	cc.
ract and the warm solution a		Citrie Acid	0.2	g.
ime to the pectin mucilage.		Aluminum Acetate	0.3	g.
he water has been added, he niform solution results, avoi		Menthol Hamamolia Water	0.5 5	g.
eating. The oil is then emul	lsified with	Hamamelis Water Perfumes (as above)		cc.
his solution, preferably in a		Alcohol (40%)	92.5	
r a homogenizer.		No. 3		
		Glycerin	1	cc.
Bathing Milk		Alum	1	g.
Emulsion of:		Zinc Sulphophenylate	0.5	
Turkey Red Oil Neutral-		Perfumes (as above) Menthol	0.5	cc.
ized with Caustic Potash	200 g.	Isopropyl Alcohol	10	cc.
Perfume Mixture	350 g.	Rose Water	10	cc.
Add then:		Alcohol (30%)	76.5	cc.
Potassium Carbonate So- lution (20° Bé.)	50 g.	And the second section of the s		
Clear Liquid Soap (10%)	400 g.	Eau de Quinine	200	
A higher content of etheric of		Alcohol Water	600 400	g.
ates more turkey red oil a	nd potash,	Quinine Sulphate	5	g.
nd eventually terpineol.	1 100	Saponine	1	g.
For a thicker balm: Use of Furkey Red, but add 100-15	mly 100 g.	Saffron Tincture	2	ğ. 2 g.
acid, and saponify the whole	with caus-	Orseille (Red Dye)	0.2	2 g.
ic.	True Strub	Rose Oil Musk, Tincture	$\frac{2}{1}$	g.
The milky character is bette		Lemon Oil	i	g.
lition of potassium stearate,	triethanol-	WAT THE PROPERTY OF THE PROPER		O
amine stearate (or oleate).		Eau de Cologne (500	%)	
		Bergamot Oil	10	cc.
Benzoin Milk		Lemon Oil	14	
Mix in a mortar or dish:		Citral	$\frac{1.4}{2.6}$	
a. Tincture of Benzoin Alcohol (95%)	50 cc.	Thyme Oil, White Rosemary Oil	3.4	
" (Alcohol (95%)	200 cc.	Lavender Oil	10	
b. Glycerin	100 cc.	Ixolene, Extra		cc.
c. Water, Distilled	700 cc.	Alcohol	500	cc.
First grind a, add b, and b		Water	500	cc.
under stirring c into a and b . a week. Filter. Shake before		Chypre Head Lotic	n.	
a moder a receive mand notice		Geraniol, C.P.	1.4	ec.
C1 . W .11 .1 TYT	ter	Cedar Wood Oil, Rectified	0.25	
(Historian Collette Met		Benzyl Acetate, Chlorine-		
Glycerin Toilette Wat	50 ~	Denzyl Acetate, Ontolline	×	
Alcohol (95%)	50 g.	Free	0.6	cc.
	50 g. 0.4 g. 50 g.		0.6	cc.

Storax Oil	0.25	cc.
Geranium Oil, Réunion	0.6	cc.
Benzyl Benzoate	2.5	cc.
Linalyl Acetate	0.8	cc.
Linalool, Extra	1.2	cc.
Anise Aldehyde	0.1	cc.
Iris Oil, Genuine, Concrete	e 0.05	cc.
Coumarin	0.15	g.
Civet, Genuine (100%)	0.02	g.
Patchouli Oil, Genuine	0.2	čc.
Musk, Artificial,		
"Ambrette"	0.2	g.
Musk, Artificial,		_
"Ketone"	0.05	g.
Labdanum Extract	0.15	cc.
Vanillin	0.13	g.
Phenylethyl Alcohol	0.6	cc.
Rosemary Oil	0.05	cc.
Alcohol	670	cc.
Distilled Water	320	cc.

Alcoholic Sulphur Hair Lotion

Sulphur Glycerin Solution	
(24%)	5 g.
Water	20 cc.
Salicylic Acid	0.5 g.
Menthol	0.3 g.
Alcohol (24%)	70 ec.
Perfume	to suit

Preparation for Head Massage German Patent 616,362

Lauryl Sulphonate	25 g.
Buckwheat Flour	30 g.
Henna	10 g.
Salicylic Acid	5 g.
Sulphur	5 g.
Castor Oil	5 cc

Scalp Stimulant

200001		
Deodorized Kerosene	80	oz.
Resorcinol Monoacetate	3	oz.
Lanolin	10	oz.
Diglycol Laurate	7	oz.

Hair Wave Concentrate

Karaya Gum	3 g.
Glycol Bori-Borate (Liquid) 6 g.
Rub together until smooth.	Stir in
Alcohol, Anhydrous	48 g.

Hair Setting Concentrate

Karaya Gum	12 g.
Glycerin or Glycol	12 g.
Alcohol	30 cc.
Perfume	to suit

The above is added to one pint of water for use.

Liquid Hair Fixative

Tragacanth, Powder	0.2 - 0.5	g.
Glycerin, C.P.	5-10	g.
Alcohol (95%)	1	g.
Distilled Water	93.8-88.5	ec.

Dissolve gum in hot water, adding it together with the glycerin (ground together previously), filter; perfume with water soluble essential oils, or use orange flower (rose flower) water instead of distilled water, then dye pale green.

If paste is wanted for collapsible tubes, use 3-4 g. of gum tragacanth.

Brilliantine

27112210110110		
Oil of Bitter Almond	1.5	ec.
Oil of Clove	3	ce.
Oil of Bergamot	6	cc.
Castor Oil	50	cc.
Clyceryl Monoricinoleate	50	g.
Suet	50	g.

Non-Greasy Brilliantine

Diglycol Laurate	40 cc.
Alcohol	60 ec.
Perfume and Color	to suit

Hair Fixative Creams

The simplest type of fixative cream is a tragacanth mucilage containing up to 25% of liquid paraffin, more or less emulsified. Such creams require vigorous shaking, as the oil separates on standing. Permanent creams which now enjoy tremendous popularity, thanks to good advertising and their own inherent good qualities, are of two types:—oil-inwater emulsions and water-in-oil emulsions, the oil in both cases being mainly liquid paraffin. The most popular of these new fixatives is of the second type, a water-in-oil emulsion. It is not, as it is often supposed, a triethanolamine emulsion, but resembles a semi-liquid cold cream. A formula for this type of cream, which has been published and widely quoted, is as follows:

Formula No. 1

Liquid Paraffin	3000 ec.
White Beeswax	100 g.
Borax	6 g.
Water	150 cc.
No. 2	
Liquid Paraffin	45 ec.
Stearic Acid	5 g.
Water	49 cc.
Triethanolamine	1 cc.
Perfume	to suit

Add the liquid parafin and stearin heated to about 65° C. to the solution of triethanolamine in water at the same temperature, and stir until it thickens. When nearly cold add the perfume. Avoid too vigorous stirring which causes

This formula gives a very thick cream which can easily be thinned by diluting

with water if desired.

Hair Fixative Perfumes

The popular ingredients include the citrus oils (orange, lemon, bergamot and lime), lavender, rosemary, geranium, petitgrain and coumarin; about 1% of perfume is sufficient. The following table will serve as a guide:

Formula	No. 1	No. 2	No. 3	No. 4	No. 5	No. 6
Bergamot Oil Lavender Oil Lemon Oil Orange Oil Lime Oil Petitgrain Oil Rosemary Oil Geranium Oil Coumarin	55 cc. 10 cc. 3 cc. 5 cc. 5 cc. 15 cc. 2 cc.	20 cc. 50 cc. ———————————————————————————————————	45 cc. 20 cc. 5 cc. 5 cc. 25 cc.	40 cc. 50 cc. — — — — — 10 g.	50 cc. 	40 ec. 40 ec. ————————————————————————————————————

Hair Oil

Formula No. 1

Alcohol, Ethyl	400 cc.
Glycerol	200 cc.
Perfume, as desired, about	1 cc.
Salicylic Acid	2 g.
Water	400 cc.
No. 2	
Alcohol, Ethyl	400 cc.
Glycerol	300 cc.
Perfume, as desired, about	1 cc.
Salicylic Acid	2 g.
Water	300 cc.

Lavender coloration may be effected by the addition of traces of ferric chloride. The preparation is completely water soluble, hence readily removed by washing, yet it serves as an excellent "stay-comb."

Soapless Shampoo

Lohrino	I (Wetting	Out			
Agent	i) `			450	g.
Mineral	Oil			50	ğ.
Alcohol				300	g.
Water		- 1	to	make 1	ĭ.

Soapless Shampoo Powder

Borax	25	oz.
Sodium Bicarbonate	25	oz.
Soda Ash	48	oz.
Saponin	2	oz.
-		

"Oil-Hair Wash" Formula No. 1

Diethylaminoethyloleyl Cit-		
rate	15	g.
Chamomile Extract	1	cc.
Lemon Juice	2	cc.

Water, Distilled, or Alcohol (50%) No. 2	}	81.5	cc.
NO. 2			

Rape Seed Oil	50	cc.
Hazelnut Oil	30	cc.
Spike Lavender Oil	5	cc.

Egg Shampoo

Prepare just before use.

Separate the yolks and whites of four or more eggs in separate bowls. To the yolks add a tablespoonful of cold water and beat until uniform with an eggbeater. Wash off the beater and beat the whites until fluffy and firm. Add the beaten yolks to the whites and fold the former into the latter. The hair is washed and rinsed with lukewarm water. Then work the egg shampoo, a little at a time, into the scalp and hair. Finally wash and rinse the hair with a strong spray of tepid (not hot) water.

Shampoo Powder

Sulphonated Lorol or		
Lohrinol	40	g.
Borax	40	ğ.
Sodium Sesquicarbonate	20	g.
This gives an excellent lather		-

Many such additions will suggest themselves to those who wish to experiment. Some people include a specially prepared saponin, 2 to 5%, to help the lather-producing properties.

Liquid Hair Shampoo

Potash Soft Soap	50	g.
Potassium Carbonate	5	g.

Glycerin 7 g. Benzaldehyde 0.25 g. Distilled Water 938 cc.

The procedure is to dissolve the soft soap, with gentle heating, in half the water. The potash, glycerin and benzaldehyde are incorporated in the rest of the water. After the two solutions have been well mixed by stirring, the finished product is left for a week before decanting, filtering and bottling. At first the perfume will be found to disappear, owing to the splitting up of the benzaldehyde into sodium benzoate and benzyl alcohol—but after the lapse of some days the characteristic almond odor will reappear, owing to the oxidation of the alcohol back to the aldehyde.

In the above formula, the soap content may naturally be increased if desired—also a proportion of alcohol may be added. Instead of the almond perfume imparted above, a stable fougère or similar compound can be employed. Likewise pine tar, or a 10% solution of henna, may be incorporated in the case of antiseptic or liquid henna shampoos respectively. Novel ingredients for imparting a pleasantly "medicated" odor

include iso-thymol.

In the manufacture of liquid soap shampoos, careful control at all points is essential. Turbidity must at all costs be avoided, and for this reason distilled water only should be used and the soap itself completely saponified. Unless proper facilities are available for saponification on the premises, it is better to purchase a ready-made soft soap base (carefully standardized examples of which are now on the market).

Shampoos should, in certain cases, be aged for even longer than a week (e.g., 15 to 30 days), then decanted into a tank fitted with a refrigerating coil, chilled to a low temperature and finally filtered through asbestos. It has been suggested that the period of aging can be radically reduced by first running the shampoo through a colloid mill or homog-

enizer.

Hair Wash

Liquid Soap		90-95	oz.
Triethanolamine	Laurate	10-5	oz.
Alcohol		10-5	OZ.

Hair Washing Soaps Formula No. 1 (for Oily Scalp) Coconut Oil 11,000 g. Castor Oil 4,750 g. Caustic Potash (50%) about 7,515 g.

Distilled or	
Softened Water 76,000	cc.
Perfume, or	
Chamomile Extract, or	
Wood Tar, Pure, or Better	
Perfume Blended with	
Extract 500- 2,000	cc.
No. 2	
Coconut Oil 11,000	or.
Olive Oil 4,750	
	8.
Caustic Potash	
(50%) about 7,520	\mathbf{g} .
Distilled or Softened	
Water 76,000	
Perfume or Extract 500-2,000	cc.
No. 3 (for Dry Scalp)	
Coconut Oil 15,000	g.
Olive Oil 6,000	g.
Caustie Potash (50%) 10,200	
	g.
Alcohol (95%) 6,000	
Distilled or	
Softened Water 53,000	00
Perfume or Extract 500- 2,000	
refluine of Extract 500-2,000	CG.

Dandruff Remover

Mercury Bichloride	0.5	g.
Resorcinol	5	g.
Alcohol	125	cc.
Water	125	ec.

Dissolve the bichloride and the resorcinol in the water. Then add alcohol. Apply on the dry scalp and rub thoroughly—then shampoo the hair. One treatment a week is usually sufficient for a complete absence of dandruff.

Dandruff Lotion

2	oz.
4	oz.
10	oz.
1	oz.
1	oz.
0.5	oz.
82	oz.
	10 1 1 0.5

Henna, White

Henna white is a bleach, varying in composition with various users. One formula, sodium perborate, 18 g.; henna leaves, 2 g.; affords an excuse for the name. No other excuse can be seen for the waste of henna leaves. Some use

Magnesium Carbonate	68 g
Sodium Perborate	32 g

Make into a paste a 50-50 mixture of hydrogen peroxide and water before use.

Birch Water

Birch Bud Oil	10	g.
Glycerin	40	g.
		70

	Soap Spirit	250	g.	Orange Flower Water,	4.00	
	Ethanol or	250		Triple	100	cc.
	Isopropyl Alcohol Bergamot Oil	650_{-5}	g.	Alcohol	800	cc.
		5	g.			
	Geranium Oil	1	g.	Eau de Lavende, Am	brée	
	Orange Flower Oil	50.0	g.	Lavender Oil, French Bergamot Oil Musk Infusion	50	cc.
	Water	50	\mathbf{g} .	Bergamot Oil	12	cc.
	***************************************			Musk Infusion	12	
	Florida Water			Ambreine	8	cc.
	Neroli Oil, "Bigarade"	5	cc.	Lemon Oil	6	
	Lavender Oil, English		cc.	Benzoin Infusion		cc.
	Bergamot Oil	30		Idola	2	cc.
	Limette Oil	2		Alcohol (96%)	2500	cc.
	Clove Oil		cc.	Water, Distilled	500	cc.
	Cassia Oil		cc.	Alcohol (96%) Water, Distilled		
	Cinnamon Oil	1	cc.	Eau de Cologne		
	Rose Oil	5	cc.			
	Ambra, Liquid, Artificial	2	cc.	Formula No. 1		
	Ambra, Liquid, Artificial Orange Flower Water, Trip	le 100	cc.	Lemon Oil	18	g.
	Alcohol (90%)	900	cc.	Bergamot Oil	16	g.
				Orange Oil, Sweet	5	g.
	777			Lavender Oil, Extra	4	g.
	Hungary Water			Mandarin Oil	$\tilde{3}.2$	g.
	Rosemary Oil	20	cc.	Petitgrain Oil, Grasse Benzoin Resinoid	3.2	œ.
	Verveine Oil	7	cc.	Benzoin Resinoid	3.2	g.
	Portugal Oil	1.5	cc.	Petitgrain Oil, Grasse Benzoin Resinoid Neroli Oil, Original Orange Oil, Bitter	2.8	g.
	Limette Oil	1	cc.	Orange Oil, Bitter	2.8	g.
	Peppermint Oil	0.5	cc.	Little Oil	4.1	g.
	Rose Water, Triple	100	cc.	Rosemary Oil	1	$\mathbf{g}.$
	Alcohol (90%)	800	cc.	Eugenol	0.6	$\mathbf{g}.$
	Rosemary Oil Verveine Oil Portugal Oil Limette Oil Peppermint Oil Rose Water, Triple Alcohol (90%) Let stand up to 6 months	before	mar-	Cumin Aldehyde (10%)	0.5	$\mathbf{g}.$
k	ceting.			Cumin Aldehyde (10%) Muscatel Sage Oil Hysop Oil	0.3	g.
	A			Hysop Oil	0.1	\mathbf{g} .
	Eau de Lubin			i taraamam tii	U. 1	g.
	Alcohol	650	ec.	Iris, Concrete (10%) Alcohol (96%)	0.1	
	Postucel Oil	1.2		Alcohol (96%)	1800	cc.
	Portugal Oil Neroli Oil	0.6		Water, Distilled	200	cc.
	Tesmina Absolute	0.6	ee.	No. 2 Bergamot Oil		
	Jasmine, Absolute Myrtle Oil	3	00.	Bergamot Oil	20	g.
	Government Oil Fronch	19	ee.	Lemon Oil	14	g.
	Geranium Oil, French Lemon Oil	3.2	ee.	Lavender Oil	5 5	g.
	Bergamot Oil	9	cc.	Benzoin Resinoid	5	g.
	Civet Tineture	3	cc.	Nerosol	5	g.
	Civet Tincture Castoreum Tincture Peruvian Balm	3	cc.	Orange Oil, Sweet	1	C)*
	Peruvian Balm	3	cc.	Mandarin Oil	4	g.
	Musk Tincture	3 3 6	cc.	Petitgrain Oil, Paraguay	2.6	g.
	Tolu Balm Tincture	6 24	cc.	Rosemary Oil	2.3	g.
	Benzoin Tincture	24	cc.	Neroli Oil	2	g.
	Marken 'l'inatiira	th.	cc.	Muscatel Sage Oil	2	g.
	Clove Tincture	60		Mandarin Oil Petitgrain Oil, Paraguay Rosemary Oil Neroli Oil Muscatel Sage Oil Jasmine Aldehyde Resinoid Iris Alcohol (96%) Water, Distilled No. 3	0.7	g.
				Resinoid Iris	0.5	g.
	The second secon			Alcohol (96%)	1800	ec.
	Aqua Mellis			Water, Distilled	200	cc.
	Honey	5	g.	No. 3		
				Lemon Oil	20	g.
	Lavender Oil, French	1	cc.	Heliotropin	7 -	g.
	Clove Oil	1	cc.	Bergamot Oil, Natural	5	g.
	Mace Oil	0.5	cc.	Bergamot Oil, Artificial Terpinyl Acetate	6	g.
	Coriander Oil	1	cc.	Terpinyl Acetate	4	g.
	Sandal Wood Oil	3.5	cc.	Neroli Oil, Artificial	4	g.
	Benzoin Resinoid	5	cc.	Orange Oil, Sweet	4	g.
	Musk Tincture (2%)	2	cc.	Coumarin	2.5	g.
	Bergamot Oil Lavender Oil, French Clove Oil Mace Oil Coriander Oil Sandal Wood Oil Benzoin Resinoid Musk Tincture (2%) Rose Water, Triple	100	cc.	Benzyl Acetate	1.5	g.

			A CONTRACTOR OF THE PROPERTY O		
Ketone Musk	0.7	g.	Eau de Cologne for th	e Bath	
Ketone Musk Citral Alcohol (96%) Water, Distilled	0.6	g.	Bergamot Oil, Free of		
Alcohol (96%)	1600	ec.	Terpenes	17	ec.
Water, Distilled	400	cc.	Petitgrain Oil, Free of Terpenes Rosemary Oil Citral	7.4	
			Terpenes	175	cc.
Ambre Eau de Col-	ogne		Rosemary OII	1.75	ee.
Reresmot Oil	20	or	Citral Tincture of Benzoin Orange Flower Water Alcohol (96%) Water, Distilled	56	cc.
Lemon Oil	20	g. 0.	Orange Flower Water	340	ee.
Heliotropin	7	g.,	Alcohol (96%)	800	cc.
Ambrette Musk	2.6	g.	Water, Distilled	3600	cc.
Lavender Oil	2.6	g.			
Petitgrain Oil, Paraguay	2.6	g.	T T T		
Methyl Ionone	2.6	$\mathbf{g}.$	Bay Oil Menthol Glycerin, C.P. Glycerin (Soap Lye) Rum Essence Alcohol (96%) Water, Distilled	8	or.
Vanillin	2	g.	Menthol	16	e.
Posamary Oil	0.7	g.	Glycerin, C.P.	16	g.
Neroli Oil	0.7	g.	Glycerin (Soap Lye)	20	g.
Coumarin	0.7	e.	Rum Essence	80	g.
Ambre, Artificial	0.6	g.	Alcohol (96%)	2000	ee.
Rose Absolute, Synthetic	0.1	g.	Water, Distilled	800	ec.
Alcohol (96%)	1800	cc.	Many managamini, asa and militalih piday managi managi militalih di Militalih Militali		
Ambre Eau de Col- Bergamot Oil Lemon Oil Heliotropin Ambrette Musk Lavender Oil Petitgrain Oil, Paraguay Methyl Ionone Vanillin Rose Oil, Artificial Rosemary Oil Neroli Oil Coumarin Ambre, Artificial Rose Absolute, Synthetic Alcohol (96%) Water, Distilled	200	cc.	Eau de Lavende	:	
			Lavender Oil, Barrême (France) Musk Infusion Ambre Infusion Bergamot Oil Lemon Oil Jasmine Aldehyde Phenyl Ethyl Alcohol Alcohol (96%) Water, Distilled Perfumes for Shaving		
Chypre, Eau de Col			(France)	40	ee.
			Musk Infusion	12	cc.
Lemon Oil	18	g.	Ambre Infusion	12	ϵc .
Lemon Oil Bergamot Oil Rose Oil, Artificial Lavender Oil Coumarin	10	g.	Bergamot Oil	12	ec.
Lavender Oil	4.	g. g.	Lemon Oil	6	cc.
Coumarin	$\frac{1}{4}$	g.	Dhami Ethil Alaskal	0.6	cc.
Sandai Wood Oil, East In	idia 2.6	ω.	Alcohol (96%)	1100	00.
Ketone Musk	2.6	g.	Water Distilled	300	cc.
Ketone Musk Vetivert Oil, Java Rosemary Oil	2	g.	77 (100,14)	0.00	
Rosemary Oil	2	g.	Perfumes for Shaving	Cronmo	
Muscatel Sage Oil, Artific Iso-Eugenol Patchouli Oil Vanillin Neroli Oil Thyme Oil	ial 2	\mathbf{g} .	C .		
Detabouli Oil	0.7	g.	Eau de Cologne Per		
Vanillin	0.7	g.	Bergamot Oil	100	g.
Neroli Oil	0.5	g.	Lemon Oil	50	g.
Thyme Oil	0.5	g.	Portugal Oil	30	g.
Mousse de Cliène, Absolut	e 0.5	σ .	Layendar Oil	20	g.
Alcohol (96%)	1800	cc.	Petitorain Oil	30	8.
Alcohol (96%) Water, Distilled	200	cc.	Bergamot Oil Lemon Oil Portugal Oil Rosemary Oil Lavender Oil Petitgrain Oil Neroli, Synthetie	20	e.
**************************************					Θ.
7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7			Bitter Almond Peri	ume	
Eau de Cologne "R	usse''				00
Lemon Oil	9	$\mathbf{g}.$	Bitter Almond Oil Bergamot Oil Lavender Oil	10	ee.
Bergamot Oil Methyl Ionone Heliotropin Lavender Oil Iso-Eugenol Vanillin	9	g.	Lavender Oil	5	ce.
Methyl lonone	6	g.			
Lexandar Oil	4		Fancy Perfume		
Tso-Eugenol	3	g.			
Vanillin	2.6	g.	Lavender Oil	150	
Ketone Musk	2	g.	Portugal Oil Bergamot Oil, Synthetic	$\frac{450}{750}$	
Rosemary Oil	$\bar{2}$	g.	Lemon Oil	150	
Linalyl Acetate	2	g.	Benzaldehyde		cc.
Ambrette, Musk	0.7	g.			00.
Neroli Oil	0.7	g.	A.T		
Coumarin	0.6	g.	Almond Perfume		
Ambre, Artificial		g.	Peru, Balsam	100	g.
Alcohol (96%) Water, Distilled	$\frac{1800}{200}$	cc.	Heliotropin Musk Tineture	125	g.
wast, Distilled	200		Musk, Tincture	96	g.

Vanillin Almond Oil Neroli, Synthetic	15 g. 10 g. 5 g.
Lavender Perfume Lavender Spike Oil Geranium Oil Coumarin Sandal Wood Oil Bergamot Oil Lemon Oil	75 g. 75 g. 75 g. 2 g. 2 g. 100 g. 25 g.
Rose Perfume Pelargol Diphenyl Oxide (1:1) Vanillin Geraniol Terpineol	100 g. 25 g. 10 g. 75 g. 20 g.
Violet Perfume Bergamot Oil Iris Resinoid Neroli Benzoin Infusion Terpineol Violet (5187, Heine) Jasmine Flower Oil Fixol—Violet	100 g. 30 g. 25 g. 75 g. 50 g. 125 g. 40 g. 50 g.
Extract, Rose Red Rose Flower Oil Nerol Phenyl Ethyl Alcohol Jasmine Aldehyde Neroli Oil Ambrette Musk Rose Absolute, Synthetic Iris, Concrete Tuberose, Artificial Bergamot Oil Narcisse, Artificial Vetivert Oil, Java Sandal Wood Oil, East India Alcohol (96%) Water	40 cc. 30 cc. 20 cc. 16 cc. 12 cc. 10 cc. 9 cc. 5 cc. 2 cc. 1 cc. 1 cc. 1500 cc.
Lilac Perfume Anisic Aldehyde Jasmine, Synthetic Heliotropin Phenyl Ethyl Alcohol Phenyl Acetaldehyde Oil Bergamot Musk Ketone Styrax Resin Oil Ylang Ylang Terpineol Individual touches may be in	10 cc. 10 cc. 5 cc. 5 cc. 5 cc. 2 cc. 2 cc. 55 cc. mparted to
Oil Ylang Ylang	2 cc. 55 cc. mparted to

Individual touches may be imparted to the above by the sparing use of any or all of the following: amyl salicylate, acetophenone, methyl anthranilate, benzyl acetate, cinnamic alcohol, benzyl benzoate, hydroxycitronellol, and oil nutmer. Perfume for Cholesterin Creams

1. Orange Flower Water instead of water:

Neroli Oil, Artificial 9 g. Aubépine 1 g. 2. Rose Water instead of distilled

water:
Rose Oil 1 g.

Geranium Oil, African 1 g.
Bergamot Oil 5 g.

3. Rose Water instead of distilled water:

 $\begin{array}{cccc} \text{Geranium Oil} & & 5 \text{ g.} \\ \text{Anisaldehyde} & & 5 \text{ g.} \\ \text{Linalylacetate} & & 2 \text{ g.} \\ \text{Eugenol} & & 1 \text{ g.} \end{array}$

The three mixtures are added to creams made with Rose Water or Orange Flower Water instead of distilled water. (Usual percentage of perfume.)

PERFUME BASES

	New Mown Hay	Chypre	Locust
Alpha Ionone Citronellol Amyl Salicylate Anisic Aldehyde Coumarin Vanillin Heliotropin Linolool Petitgrain Jasmine, Artificial Patchouli Oil Aldehyde C ₁₀ , 50% Iso Eugenol Phenyl Ethyl Alcohol Musk Xylol Copaiba, Balsam Birch Tar Lemon Oil Bergamot Oil Bergamot Oil Rose, Artificial Cedar Oil Phenyl Acetic Aldehyde, 50% Phenyl Acetic Acid Hydroxycitronellol Cinnamic Alcohol	10 20 100 20 5 5 7 10 10 20 1 1 5 ————————————————————————————————	25 	5.5
Cananga Oil	-	-	3
ate, 5%	-	,	1.0
Amyl Cinnamic Aldehyde	1	=	1.3 5

		·		
				4
	Flowery Bouquet	Bouquet	tal	tal
	M.	bn	Orienta	Orienta
	110 301	301)ri	Ï.
	Ŧ			O
Rose Geranium Oil	100			
Rose, Artificial	20			
Valley Lily, Artificial	500	500	350	100
Terpineol	200	_	110	100
Hydroxycitronellal	200			
Bois de Rose	200			
Coumarin	30	100		_
Anisic Aldehyde	20		30	30
Methyl Anthranilate.	150	20	_	_
Civet Tincture	50		100	60
Hyacinth, Artificial	100			_
Benzyl Benzoate	200	200	200	100
Musk Ambrette	50		50	30
Opoponax	10		200	100
Oak Moss, Liquid	200		100	50
Cananga Oil	100			
Lavender Oil		20	20	10
Bergamot Oil		100	_	
Cassia Oil		10		
Tuberose, Artificial .		100		
Methyl Heptine Car-			l	
bonate, 5%		100		
Geraniol		100		
Vanillin		100		_
Musk Ketone	_	50		
Orange Blossom, Ar-				
tificial			610	100
Jasmine, Artificial	-		440	
Vetivert Oil	-			100
Jasmine Aldehyde				200
Petitgrain Oil		-		100
Phenyl Ethyl Alcohol				30
Linalyl Acetate				50
Linalool			-	50

	Flowery Bouquet A	Bouquet A	Bouquet B	
Aldehyde C	20	20	20	
Oak Moss, Liquid	100	_		
Jasmine Liquid, Absolute	500	200	200	
Rose, Artificial	500	1000	300	
Iso Butyl Salicylate	200	100	100	
Methyl Ionone	500		300	
Lilac, Artificial	500	200	300	
Musk Ketone	200	_	200	
Methyl Heptine Carbonate,				
5%	50		-	
Valley Lily, Artificial	500	200		
Bois de Rose	200	200	200	
Melittis (Givaudan)	200			
Orange Blossom, Artificial	1-	300	300	

Methyl Phenyl Acetate		40 100 50 30	
Musk Ambrette		100 50	
Para Cresyl Phenylacetate		50	
Vanillin			
Vanimin		00	I
Aldohydo C 501		100	100
Aldehyde C ₁₀ , 5% Olibanum Gum, 2:1		150	200
			200
Terpineol			200
Hydroxycitronellal			100
Cananga Oil			100
Coumarin			30
Anisic Aldehyde	-		20
Methyl Anthranilate			100
Civet Tincture			50
Labdanum			100
Coriander Oil			20
Castoreum, 10%	-		100
Ambergris Tincture			100

	Chypre A	Bouquet C	Bouquet D
Jasmine, Artificial Musk Ketone Oak Moss, Liquid Bergamot Oil Rose, Absolute Patchouli Oil Musk Tincture Vanillin Coumarin Indol, 5% Hydroxycitronellal Lemon Oil, Terpeneless Phenyl Ethyl Alcohol Methyl Ionone Aldehyde C9, 50%	200 400 500 1000 400 500 200 100 200 200 30	200 100 — — — — —	80 500
Methyl Heptine Carbonate, 10%			150
Bois de Rose Cassie, Artificial Benzyl Benzoate Diethyl Anthranilate Linalyl Acetate Benzyl Acetate Tolu, Balsam		200	60 1000 50 300 300 300
Rose, Artificial	-	-	an

	French Type	French Lilac Type
Oak Moss, Liquid Bergamot Oil, Terpeneless. Linalyl Acetate Sweet Orange Oil Valley Lily, Artificial Narcissus Absolute Jasmine, Artificial Rhodinol Alcohol C9 Aldehyde C9, 5% Linalool Geranyl Acetate Methyl Phenylacetate Alpha Ionone Vetivert Oil Terpineol Coumarin Vanillin Musk Ketone Canada Snake Root Oil Hydroxycitronellal Geraniol Phenyl Acetic Aldehyde, 50% Phenyl Ethyl Alcohol Anisic Aldehyde Rose, Artificial Labdanum	200 150 50 200 300 100 400 200 70 100 200 100 100 100 100 100 100 100 10	2000

	Bouquet E	Violet
Raldeine D Lemon Oil Rhodinol Alpha Ionone Hydroxycitronellal Cananga Oil Aldehyde C ₁₂ , 5% Methyl Heptin Carbonate, 10% Cassie, Artificial Guaiac Methyl Ionone Orris, Liquid, 10%	300 20	100 1000 1000 300 100 100 200 100 300 100 175

	Bouquet E	Violet
Bergamot Oil, Terpeneless. Linalyl Acetate Jasmine, Artificial Aldehyde C ₉ , 5% Vetivert Coumarin Rose Geranium 0il Rose, Artificial Bay Oil, Terpeneless Eugenol Petitgrain Oil Bergamot Oil Indol, 5% Ambreol Lavender	200 100 500 100 100 400 200 100 400 300 150 500 150	100

	Jasmine	Sweet Pea	Heavy Oriental
Benzyl Acetate Bergamot Oil Bois de Rose Benzyl Alcohol Phenyl Ethyl Alcohol Indol, 5% Hydroxycitronellal	1500 150 150 300 300 50 250	300	3000 200 — — 200
Orange Blossom, Artificial	250 150 300 100 — —	200 200 — 150 450 50	200 — 100
50% Terpineol Iso Butyl Phenylacetate Rose, Artificial Tolu Alcohol C ₉ Benzyl Benzoate Anisic Aldehyde Lavender Oil		100 1000 120 80 150 60 150	400 100 60
Tolyl Acetate Vanillin Oak Moss, Liquid Aldehyde C ₁₀ , 5% Diethyl Anthranilate Ambreol			100 200 400 160 580 600

	Carnation	Honeysuckle
Eugenol Jasmine, Artificial Heliotropin Rose, Artificial Phenyl Ethyl Alcohol Orange Blossom, Artificial Ocillet Orris Liquid, 10% Musk Ketone Ambreol Benzyl Iso Eugenol Bergamot Oil Indol, 5% Hydroxycitronellal Benzyl Acetate Benzyl Formate Benzyl Formate Benzyl Formate Benzyl Propionate Benzyl Propionate Benzyl Pensoate Bois de Rose Aurania Dananga Oil Amyl Cinnamic Aldehyde Para Cresol, 10% Petitgrain Oil	1600 400 400 100 50 100 100 100	

	Lilac	Rose	Orange Blossom	Heavy Modern Oriental	
Citronellol Cananga Oil	10 20	30 5	3		
Amyl Cinnamic Aldehyde Methyl Acetophenone Hydroxycitronellal Phenyl Ethyl Alcohol Linalool Terpineol Methyl Para Cresol Musk Ketone Valley Lily, Artificial Iso Eugenol Aldehyde C ₁₀ , 5% Benzyl Acetate Geraniol Ionone	10 5 10 11 10 20 1 5	5 20 5 — — 5 10 50 5	10 10 10 	50	
Geranyl Acetate	=	10	_		
Copaiba Balsam Patchouli Oil	_	10		_	
Phenyl Acetic Acid	_	2 2	2		

,	Lilac	Rose	Orange Blossom	Heavy Modern Oriental
Linalyl Acetate		3		
Petitgrain Oil			100	
Methyl Anthranilate.			15	
Beta Naphthyl Ethyl				
Ester	-		10	50
Amyl Salicylate				30
Ionone				10
Benzylidene Acetone.			-	6
Musk Xylol			******	5
Vanillin				3

Lily-of-the-Valley Flower	Oil	
Geraniol, from Palmarosa Oil Linalool, from Rosewood Oil Phenylethyl Alcohol Phenylacetaldehyde Dimethyl	$\frac{12.5}{15}$	g 95 g
acetal a-Ionone Benzaldehyde	5 $ 1.5 $ $ 0.1$	9 9 9 9 9 9
Jasmine Flower Oil, Artificial Rose Oil, Artificial,	10	g.
Extra Fine Lilac Flower Oil, Artificial Ylang Ylang Oil, Manila	25 4	ಚ ಚ ಚ
Rhodinol Coriander Oil, Terpene-Free Hydroxycitronellal Dimethyl-	10 0.5	g. g.
acetal Hydroxycitronellal Diethylacetal	20 40	g. g.

Lilac Flower Oil		
Ylang Ylang Oil, Manila Jasmine Flower Oil, Arti-	1	g.
ficial	12	g.
Rhodinol	6	g.
Acacia Flower Oil, Artificial	2	g.
Hydroxycitronellal Diethyl-		
acetal	30	g.
Terpineol, Extra	20	g.
Phenylacetaldehyde		
Dimethylacetal	4	g.
Aubépine (from Anethol)	2	g.
Heliotropin	12	g.
Iso-Eugenol	1.5	g.
Vanillin	0.5	g.
Octyl Acetate (10%) in		Θ.
Benzyl Alcohol	0.5	g.

Perfume Oil, Type "To Formula No. 1	osca''		pure alcohol and kept in the ing from time to time, and fil a few weeks.		
Orange Oil, Sweet,	0.5		No. of Salah		
Calabrian	8.5	cc.	70 4 013 77 // 0 7		
Bergamot Oil, Extra Fine,	17	00	Perfume Oil, Type "Quelqu		eurs ' '
Reggio	17	ec.	Tart ("Herb") Ty	pe	
Lemon Oil	19	cc.	Formula No. 1		
Ylang Ylang, Genuine	6	cc.	Olibanum Oil	3	00
Rose Oil, Genuine,	2.5	ec.	Geraniol, C.P.	7.5	cc.
Bulgarian Jasmine, Pure	$\frac{2.3}{1.3}$	cc.	Alpha Amyl Cinnamic	1.0	CO.
Coumarin	6.5		Aldehyde	2.36	00
Musk, Artificial,	0.0	g.	Citral	5	cc.
"Ambrette"	1	g.	Geranium Oil, Réunion	3.5	cc.
Musk, Artificial,	_	ъ.	Benzyl Alcohol	10	cc.
"Ketone"	1	g.	Linalyl Acetate	7	cc.
Cedar Wood Oil, Rectified	$\bar{5.5}$	cc.	Hydroxycitronellal, C.P.		•••
Neroli Oil, Genuine	2.5	cc.	(100%)	14	cc.
Geraniol, C.P.	4	cc.	Heliotropin, Crystallized	10	g.
Phenylethyl Alcohol	1.5	cc.	Cananga Oil, Java	13	ec.
Benzoin Extract, Filtered	5	cc.	Ionone for Soaps	4	cc.
Petitgrain Oil	1.5	cc.	Methylnonyl Acetaldehyde		
Linaloë Oil, Cayenne	6	cc.	(100%)	0.14	cc.
Sandal Wood Oil,			Benzyl Acetate, Free of		
East Indian	5.5	cc.	Chlorine	6	cc.
Indol (100%)	0.07	cc.	Linaloë Oil, Cayenne	3	cc.
Iris Oil, Genuine, Concrete	1.5	cc.	Terpineol, C.P.	11	cc.
Castoreum (100%)	0.05	g.	Musk, "Ambrette,"		
Basilicum Oil	0.03		Artificial	0.5	g.
Undecyl Aldehyde (100%)	0.05	cc.	No. 2		-
Mousse de Chêne, Liquid	0.5	cc.	Benzoin, Extract	3	cc.
Vanillin	3	g.	Olibanum Oil	1.36	
Menthol	0.5	g.	Citronella Oil, Colombo	3	cc.
No. 2			Cananga Oil, Java	10	cc.
Bergamot Oil, Extra Fine	11	cc.	Heliotropin, Crystallized Linaloë Oil, Cayenne	6	g.
Lemon Oil	26.5	cc.	Linaloë Oil, Cayenne	7	cc.
Orange Flower Water Oil,			Hydroxycitronellal, C.P.		
Genuine	1	cc.	(100%)	7	cc.
Ylang Ylang Oil, Genuine	9	cc.	Benzyl Acetate, Chlorine-		
Sandal Wood Oil,			Free	3	cc.
East Indian	8	cc.	Terpineol, C.P.	26.5	cc.
Amyl Salicylate	3.5	cc.	Citral	3	cc.
Iris Oil, Genuine, Concrete	1	cc.	Methylnonyl Acetaldehyde		
Civet, Genuine (100%)	0.22	cc.	(100%)	0.14	
Patchouli Oil	1.5	cc.	Geranium Oil, Réunion	5.5	cc.
Coumarin	4	\mathbf{g} .	Ionone for Soaps	5.5	cc.
Vanillin	5	g.	Phenylethyl Alcohol	5	cc.
Rose Oil, Bulgarian	3.5	cc.	Linalyl Acetate	4.5	cc.
Petitgrain Oil	1.5	cc.	Anise Aldehyde	6.5	cc.
Musk, Artificial,			Alpha Amyleinnamic	9	
"Ketone"	6	g.	Aldehyde	3	ec.
Geraniol, C.P.	6.5	cc.			
Benzoin Extract, Filtered	5	ec.	Parfyra Oil Type ((Ouelan	ing Till	
Undecyl Aldehyde (100%)	0.2	g.	Perfume Oil, Type "Quelqu		surs
Birch Tar Oil,	0.09		For Fine Soaps (Soft	Type)	
Twice Rectified	$\frac{0.03}{2}$		Cananga Oil, Java	9	cc.
Cedar Wood Oil, Rectified		cc.	Benzyl Acetate, Free of		
Neroli Oil, Genuine	0.5	cc.	Chlorine	5	cc.
Linaloë Oil, Cayenne	$\frac{2}{0.05}$	cc.	Ionone for Soaps	5	cc.
Opoponax Extract Jasmine Oil, Pure	2	cc.	Linalyl Acetate	6	cc.
The second secon			Linaloë Oil, Cayenne	2.3	cc.
The above-mentioned perfu	me coi	mposi-	Heliotropin, Crystallized	8	g.
tions should be made up 1-29	70 In 8	90%	Geraniol, C.P.	8	cc.

	COSME	TITOS
Musk, "Ambrette,"		
Artificial	3.5	g.
Bergamot Oil	_	cc.
Phenylethyl Alcohol		cc.
Pangral Alaskal	_	
Benzyl Alcohol	9	cc.
Alpha Amylcinnamic		
Aldehyde		cc.
Terpineol, C.P.	21	cc.
Indol, Crystallized	0.06	g.
Lemon Oil, Genuine	4.	cc.
Anise Aldehyde	$\tilde{4}$	cc.
Hydroxycitronellal, C.P.	9	
Mathedramy Acatellahyd		cc.
Methylnonyl Acetaldehyde	0.74	
(100%)	0.14	cc.
Perfume Oils "Chypre	Extract	,,
Formula No. 1		
Bergamot Oil	33	cc.
Geranium Oil, Réunion	2	
Page Oil Convine Dulmer		cc.
Rose Oil, Genuine, Bulgar	man 5.5	cc.
Ylang Ylang Oil, Genuine		cc.
Rosemary Oil	4	cc.
Coumarin	8	g.
Lavender Oil, Genuine	6	cc.
Jasmine, C.P.	2.4	cc.
Vanillin	3	
Amino Aldolando		g.
Anise Aldehyde Cedar Wood Oil, Rectifie	5.5	cc.
Cedar Wood Oil, Rectifie	d 1.5	cc.
Patchouli Oil, Genuine	0.5	cc.
Mousse de Chêne, Decolor	ized 3	cc.
Mousse de Chêne, Decolor Opoponax Extract	2	cc.
Linaloë Oil, Cayenne	18	cc.
Civet Genuina (100%)	0.6	
Civet, Genuine (100%) Musk, "Ambrette," Artif	301.15	g.
Musk, Ambrette, Arth	16191 4.0	g.
No. 2		
Lemon Oil	12	cc.
	9	
Bergamot Oil		cc.
Benzyl Acetate, Free from		
Chlorine	- 8	cc.
Cedar Wood Oil, Rectified	1 9.5	cc.
Benzyl Benzoate	6	cc.
Hydroxycitronellal, Pure		
(100%)	5	cc.
Geraniol, C.P.	7	cc.
Vanillin	4	
Danasia Fratus et Tiltone		g.
Benzoin Extract, Filtered	I 5.5	cc.
Sandal Wood Oil,		
East Indian	5	cc.
Geranium Oil, Réunion	3	cc.
Coumarin	2	g.
Rose Oil, Genuine, Bulga		cc.
Linelas Oil Coverne	95	
Linaloë Oil, Cayenne	2.5	cc.
Musk, "Ambrette,"		
Artificial	1.5	g.
Patchouli Oil, Genuine	1.5	cc.
Labdanum Extract	3	cc.
Civet, Genuine	0.3	g.
Olibanum Extract	0.7	cc.
Tria Oil Gannina Concret		cc.
Iris Oil, Genuine, Concret	ino I	
Mousse de Chene, Decolor	rized 2	cc.
Mousse de Chêne, Decolor Ylang Ylang Oil, Genuin	e 5	cc.
Phenylethyl Alcohol	5.5	cc.

Cuticle Remover

Glycerol	20	oz.
Potassium Hydroxide	4	oz.
Water	76	oz.
Perfume	0.3	OZ.
Basic Red Dye	tr	ace

The potassium hydroxide is dissolved in the water and the glycerol then added. The perfume usually used is a terpeneless lemon oil. Just enough dye is added to give same a pink color in the bottle.

Cuticle Softener Formula No. 1

Light Turbine Oil—color and perfume to suit.

No. 2		
Diglycol Laurate	10	oz.
Deodorized Kerosene	10	oz.
Perfume	to	suit
No. 3		
Olive Oil	88	oz.
Petroleum Jelly	12	OZ.
Red Dye Oil Soluble		

to a pink color trace Perfume Lilac, enough, about 0.3 oz.

A lower priced product may be prepared by using a medium bodied white mineral oil. The petroleum jelly should be nearly white. This jelly is melted at a low heat and added to the olive oil. The dye is mascerated with a small portion of the oil and this paste is used to tint the entire mass. The perfume is added in amount varying with the strength of the particular product used.

Nail Polish

rormunt No. 1	
Amyl Acetate	700 g.
Methyl Alcohol	300 g.
Nitrocellulose	50 g.
Benzoin	100 g.
Carmoisine (1% Alcoholic	100 g.
	F0
Solution)	50 cc.
*	or to suit
No. 2	
Butyl Acetate	250 g.
Ethyl Acetate	150 g.
Ethyl Alcohol	400 g.
Butyl Alcohol	200 g.
Damar	200 g.
Color	5 g.
	to suit
No. 3	
Methyl Ethyl Ketone	650 g.
Resorcinol Diacetate	100 g.
Ethyl Lactate	200 g.
Nitrocellulose	100 8.
*1. Date Cartillope	100 g.

Sandarac	5 g.
Color	to suit

Sometimes the polish is perfumed with a little ionone or ylang ylang oil, but more often this is not done.

No. 4

Nitrocellulose	
Viscosity)	225 g.
Damar	75 g.
Butyl Acetate	25 g.
Butyl Alcohol	20 g.
Ethyl Acetate	15 g.
Alcohol	40 g.
Carmine Red	sufficient to color

Nail Polish Powder

Putty Powder (Tin Oxide)	40 oz.
Infusorial Earth (325 Mesh)	55 oz.
Stearic Acid (Powdered)	5 oz.
Color (Pigment)	to suit
Perfume	to suit

Removers, Nail Polish Formula No. 1

The nail polish remover consists chiefly of the solvent alone. It has been found, however, that butyl stearate has a particularly rapid action on the film, and many modern removers make use of it in conjunction with other solvents. An effective remover can be made by mixing butyl stearate 1 part, amyl acetate 3 parts, and acetone 4 parts. Diglycol laurate is also included to prevent brittleness of nails (about 1-2%).

	No. 2	
Amyl Acetate		1 oz.
Acetone		1 oz.
	No. 3	
Amyl Acetate		1 oz.
Alcohol		1 oz.
Acetone		1 oz.
Diglycol Laur	ate	1/8 oz.

Eyebrow Pencils

Apart from those methods which serve to preserve the eye region in good physical condition, actual beauty treatment is now practiced on a very considerable scale. Coloring of the eyebrows, painting of the eyelashes and shading of the eyelids are now important components of face cosmetics, the greatest attention being devoted to the first operation. Coloring of the eyebrows or their simulation after complete shaving is effected with colored wax pencils. As already mentioned, ordinary pure charcoal pencils tend to cause falling-out and drying of the hair.

Ingredients used in preparing the wax pencils are white wax, benzoated tallow, cocoa butter, petroleum oil and olive oil. The pigments are lamp black, umber, and ochre. Large manufacturers find it economical to use pigment grinding machines and other equipment of the most modern design, but small concerns can nevertheless cope with the production of these cosmetics. The base comprises a composition made up from 110 g. fine petroleum oil, 60 g. white ceresine, 15 g. white wax, 240 g. benzoated tallow, and 1 g. coumarin. The fatty base is thoroughly ground with the pigments, the molten base being gradually stirred into the very finely powdered pigment contained in a mortar. After thorough trituration the mixture is again warmed, digested for about half an hour on a water bath, and again allowed to cool. As soon as the mass begins to thicken, it is again vigorously stirred and forced through a fine-mesh sieve by applying powerful pressure with the pestle. Lumps and impurities are retained upon the sieve. The preparation which passes through the mesh is then again thoroughly mixed, with gentle heating before casting. The mass should be neither too hot nor too fluid when being cast, since settlement of the insoluble pigment will result in lack of uniform coloration. Oilsoluble dyestuffs will certainly only enter into consideration in exceptional cases. According to another process, the melt is prepared from 2 parts cocoa butter, 2 parts ceresine, and 1 part olive oil. Into this is stirred 0.6 part dyestuffs (i.e., about 10% of the total gross weight), which has previously been ground up with a little olive oil.

As soon as the mass has reached the state when it can just be cast, it is emptied into metal moulds. As a rule these impart the required taper to the pencils, but if this is not the case they are tapered after removing from the moulds and wrapped in thick metal foil while leaving the points exposed.

Eyelid Pencils

The production of shading tones on eyelids can be effected with pencils, the composition of which is very similar to that of the eyebrow pencils. The mass consists of the fatty base detailed above with the addition of about 20% ceresine. The color scale is somewhat more varied in the case of these pencils, since a wider range of tones can be induced in the usual brown and bluish black shades. Chestnut is obtained by mixing 225 g.

pale umber and 150 g. mahogany brown with 1000 g. of the molten wax mass. For dark brown tones mix with the same quantity of wax 300 g. of a brun foncé; black shades require for the same wax quantity 100 g. zine white, 120 g. ultramarine, and 4 g. lamp black.

Regarding the perfuming of these preparations, these should generally be of a very refined character. About 5 to 10 g. of perfume are required for each kilogram of mass. In cases where a fancy perfume is desired, preference should be given to one with a fresh natural odor.

Brown Eyebrow Pencil

Burnt Sienna	80 g.
Burnt Umber	100 g.
Hard Paraffin	420 g.
Soft Paraffin, Yellow	400 g.

Eyebrow and Eyelash Softener Formula No. 1

20

OZ.

Castor Oil

Almond Oil	60 oz.
Perfume	3/4 oz.
No. 2	
Diglycol Laurate	100 oz.
Acetic Acid, Glacial	1/4 oz.
Mineral Oil, Medicinal	200 oz.
No. 3	

Beeswax	200 g.
Cocoa Butter	300 g
Melt together and add:	-
Peanut Oil	750 g.
Moldex or Other Good	J
Preservative	2 g

Lipsticks (and Eyebrow Pencils)

Paramn	22	oz.
Vaseline Oil, White	3	oz.
Beeswax, White	1	oz.
Ozokerite Ceresine	3	oz.
Titanium Dioxide	1	oz.
Colors: For 100 parts use:		
Fixation Red (Fixierrot)		

I No. 46 3.5 oz.

Medium Red (Mittelrot)

No. 28 22 oz.

Other red dyes used: Carmine, Nakarat, Fixierrot, Cherry Red, Orient Red.

After Shave Lotions Formula No. 1

Glycerin Lactic, Citric, or Phos-	2	g.
phoric Acid Menthol	$0.2 \\ 0.5$	

Alum	0.3	
Perfume	$0.5 \\ 96.5$	
Alcohol (45%)	90.0	g.
No. 2	_	
Glycerin	5	g.
Alum	1	g.
Zinc Sulphophenolate	0.5	g.
Propyl Alcohol, C.P.	10	g.
Rose Water	$\frac{10}{0.5}$	g.
Perfume	72.5	g.
Alcohol (45%)	ال • شا	g.
No. 3		
Alcohol (40%)	1000	
Glycerin, C.P.	40	g.
Aluminum Lactate	3	g.
Citric Acid	2	g.
No. 4		
Zinc Sulyhophenolate		g.
Alcohol (96%)		cc.
Witch Hazel	10	g.
Peruvian Balm	0.25	
Glycerin, C.P.	1	g.
No. 5		
Distilled Water	20	cc.
Isopropyl Alcohol	4	cc.
Alcohol	4	ec.
Alum		g.
Glycerin		cc.
Zinc Sulphophenolate	0.25	g.
No. 6 (Cloudy)		
Emulsone B	50	g.
Boric Acid	50	
Isopropyl Alcohol	100	
Diethylene Glycol	200	g.
Titanium Dioxide	60	g.
Distilled Water	4	1.
Menthol	2	g.
Moldex or Other Preservat	ive 2	g.
		-

Shaving Creams, Foaming Formula, No. 1

TOTALUM TION T	
a. Stearin Coconut Oil Fatty Acid	25 g.
". Coconut Oil Fatty Acid	8 g.
b. Caustic Potash (50° Bé.) Water Glycerin	15 g.
b. \ Water	50 ec.
{Glycerin	4 g.
c. Stearin	3 ℃.

Melt up a, then introduce the solution b with stirring. Stir until cooled, then introduce c. When homogeneous, cover container and let stand over night. Perfume is added the next morning, optionally together with alcohol. Keep 8-14 days in earthenware jars, stir with a wooden rod on each day. In this time, the cream should become softer. If not, treat with a little caustic potash solution (20° B6).

Perfume: Lavender, Rose, Violet,

90	THE	CHEMICA	L FORMULARY	
Benzaldehyde, o Chypre.	No. 2	Cologne or	$a.\begin{cases} \text{Pig Fat} & \text{No. 4} \\ \text{Olive Oil} \\ \text{Tallow} \end{cases}$	80 g 100 g. 75 g.
Bleached Olive Oil Fat: Coconut Oil F Water Caustic Potasl	ty Acid atty Acid	25 g. 25 g. 10 g. 35 cc. 25 g.	Coconut Oil Caustic Potash (38° Bé.) Glycerin Water c. Stearin	60 g.
Method as in Stearin Coconut Oil, o Olive Oil, or I Caustic Potas	No. 1. No. 3 r Fatty Acid	30 g. 15 g. 10 g. 27 g. 32 cc.	As in No. 1. Brushless Shaving Creat 1. Glycosterin	
Water Glycerin Stearin Method as in	No. 1. No. 4	6 g. 3 g.	Peanut Oil Water Moldex or Other Good Preservative	5 oz. 60 oz. 0.2 oz.
Stearin Coconut Oil Caustic Potasi Water Glycerin Turkey Red C	ı (50° Bé.)	30 g. 11 g. 17 g. 30 cc. 10 g. 2 g. e alkali	Olive Oil Lanolin Glycerin Triethanolamine Sodium Carbonate	20 oz. 6 oz. 2 oz. 6 oz. 2 oz. 1 oz. 63 oz. to suit
For	Cream, Foamir mula No. 1		Sacrilar Chair Danie	.
Coconut Oi	, or Fatty Acid I, or Fatty Acid tash (38° Bé.)	l 14 g. 28 g.	Soapless Shaving Prepara German Patent 604,77 Formula No. 1 Glycol Stearate	
b. Water Glycerin (2 c. Stearin		20 g. 12 g. 5 g.	Water No. 2 Absorption Base (Parachol)	400 g. 100 g.
Mix a in the points (lowest f. C., then stir in until cool, add	b, warm to 65°	to 60–70° C. Stir	White Beeswax Water No. 3	25 g. 100 g.
oughly, let standing stir up thor Cover, let stand, jars on next da	over night. Noughly, adding and fill into early.	ext morn- perfume.	Glycol Palmitate Petrolatum Water No. 4	100 g. 100 g. 200 g.
Bleached Paln Olive Oil Fat Coconut Oil F Water		d 50 g. 50 g. 20 g. 70 g.	Diglycol Laurate Lanolin Petrolatum Water No. 5	100 g. 100 g. 50 g. 100 g.
Caustic Potasl Method as in		50 g.	Stearic Anilide Glycol Stearate Absorption Base Water	100 g. 300 g. 100 g. 1500 g.
a. Stearin Coconut Oi Caustic Po Glycerin Water c. Stearin As in No. 1.	tash (50° Bé.)	90 g. 10 g. 42 g. 20 g. 100 g. 10 g.	No. 6 Glycol Stearate Absorption Base (Parachol) White Beeswax Sesame Oil Water Saponine	30 g. 100 g. 30 g. 800 g. 600 g. 16 g.

Shaving Creams, Non-I	Poaming
Formula No. 1 (For Fa-	tty Skin)
Stearin	50 g.
$a. \left\{ egin{aligned} ext{Stearin} \ ext{Vaseline} \end{aligned} ight.$	10 g.
$b. \begin{cases} ext{Triethanolamine} \\ ext{Borax} \\ ext{Water} \end{cases}$	1.5 g.
$b. \{ Borax \}$	1.5 g.
Water	130 cc.
c. Alcohol (Perfume)	3 g.
	~ ~

Pour a, 70° C., into b, 60° C. Cool stirring; add c before solidification. Pack in collapsible tubes.

No. 2	
Stearin	45 g.
Triethanolamine	2.5 g.
Glycerin	15 g.
Water	67.5 cc.
Witch Hazel	50 cc.
Method as in No. 1.	

Latherless Shaving Cream U. S. Patent 1,991,501

A neutral shaving preparation of a latherless type which consists of a mixture of the following ingredients in substantially the proportion stated, stearic acid 11 g., lanolin 10 g., coconut oil 0.3 g., concentrated ammonium hydroxide 1.35 g., paraffin wax 6 g., spermaceti wax 2 g., boric acid 1.5 g., water 75 g., and having a trace each of menthol, camphor and perfume.

Stearic acid and hydrous lanolin containing 20% water, together with coconut oil are melted together, and to this mixture is added the concentrated ammonium hydroxide, which contains ap-

proximately 25% of ammonia.

The waxes are then added and heating is continued until the entire mixture is liquefied. The resulting mixture is subsequently removed from the heat and a warm solution of the boric acid in approximately 75 g. of water is added while continuously stirring.

At this point, or at any point previously, the menthol, camphor and selected perfumes are added in amounts which

give the most pleasing effect.

The mixture is then violently stirred until cold, and the final resulting product is a white cream.

Shaving	Creams,	Non	-Fo	amin	g
	Formula				_

	CATTILITIES TAGE	.1.	
$a.\begin{cases} Stearin \\ Vaseline \end{cases}$		-75 g.	
Vaseline		13 g.	
Triethand	lamine	2 g.	
b. Borax Water		2 g.	
		195 g.	
c. Alcohol		6 g.	

Melt up a to 70° C., mix b and heat up to 60° C., then pour a into b with stirring. Shortly before the cooling (solidification) add perfume in the alcohol c, stir until cold. Fill into collapsible tubes.

No. 2	
Stearin	36 g.
Aminostearin	10 g.
Vaseline	5 g.
Glycerin	5 g.
Water	130 g.
No. 3	
Stearin	30 g. 10 g.
Triethanolamine	10 g.
Witch Hazel	100 g.
Water	45 g.
Glycerin	10 g.

Camphor Shaving	Milk	
Camphor, Spirits of	50	g.
Glycerin	50	g.
Lavender Oil	2	g.
Alcohol	600	g.
Add:		
Borax, Powder	25	g.
Distilled Water	1200	g.
Fresh Lemon Juice	200	g.
Stir: allow to stand over	night: f	ilter

Milky-White Shaving Soap,	Liqui	d
Coconut Oil	30	g.
Tallow	90	g.
Stearic Acid	90	
	it 90	g.
Potassium Carbonate	1	g.
Distilled or Softened Water	370	g.
Glycerin		g.
Alcohol		g.
Perfume 2	.5-10	g.

Shaving Milks Formula No. 1

roimma ivo. 1		
Mix in warmed mortar:		
Wool Fat	10	œ.
Borax	2	
Glycerin	15	œ.
Orange Flower Water	40	g.
Rose Water	40	g.
Tincture of Benzoin	10	g.
No. 2		
Make up emulsion of:		
Almond Oil	20	g.
Glycerin	20	g.
Gum Arabic	20	g
Rose Water	440	g.
And add:		

40 g.

10 g.

Glycerin

Perfume

Tincture of Benzoin

V.	5 CIIIIIII						· · · · · · · · · · · · · · · · · · ·
No. 3		Caustic	Potash (5	10%)			
Grind:				a	bout	6.33	g.
Lanolin, Pure, Pale	50 g.	Distilled	Water (or			•
Coconut Oil	25 g.		ied Water		7	9	g.
Borax	8 g.				_		0
	25 g.	_				-	
Neutral Soap Powder	20 g.	2	having S	oap,	Lıquı	a.	
Water	80 g.	Olein, L	ight			9	g.
Rose Water	400 cc.	Coconut	Oil, Cock	$_{ m in}$		3	g.
Orange Flower Water	100	Caustic	Potash (50° B	sé.)	5.3	3 g.
(Tepid)	400 cc.	Alcohol			•	1	g.
Peppermint Oil	2 ec.	Glycerin	, C.P.			8	g.
		Water				73	g.
Astringent After Shavi	ing Milk	Rose Wa	iter			1	g.
•	_						_
Formula No. 1	1	Shaving	Soap, Sin	ailar t	o ''R	asibl	loe''
Glyceryl Monostearate	10 g.	Stear				100	g.
Vegetable Oils	8 g.						-
White Paraffin Oil, Odorle	ess 2 g.	Glyce	:	4000	~~.	5	g.
Distilled Water	73 g.	h Caust	ic Potash	. (39°	Bé.)	40.	2 g.
Acetic Acid (50%)	5 g.	Caust	ic Soda (37° B	é.)	11.	4 g.
Glycerin (28° Bé.)	2 g.		ut Soap		•	30	g.
Add perfume resistant to			ach porti		d mi		
-	acrasi	in above o		iore ar	iu 1111	A to	gemer
No. 2	Ì	111 00000	ruci.	•			
Camphor	2 g.		B1000000000000000000000000000000000000				
Eau de Cologne Oil	4 g.		After Sh	ave L	otion		
Alcohol	300 g.	Alcohol	(95%)			68	0 g.
		Perfume	Oil			00	6 °g.
Glycerin (28° Bé.)	80 g.	Glycerin				1	5 g.
Rose Water	614 g.	Tannic					E
-			Water			29	4 or
Transparent Liquid Sha	ving Soap						
Olain Clear Pala	125 ~		alcohol				
Transparent Liquid Sha Olein, Clear, Pale Coconut Oil	15.5 g.	glycerin, t	nen the v	vater-t	annic	acı	1 solu-
Coconav On	1.010 g.	tion.					
POW	DERED HAN	D TOILET	SOAPS				
		No. 1	N- 0	TAT.	. 0	'nT	. 1
		No. 1	No. 2		o. 3		0.4
Formula:		Bathroom					
		Traveland				r	
		HomeUse	rage Us	e Ge	neral		
Dry Yellow Powdered Soa	p, 92% plus						
c.p.s.,* S.N.† to be over	210 titre,‡ 25						
to 35° C		75 lb.		40	lb.	60	lb.
Cocoanut soap-powder, 309	% Anhydrous						
Soap Contents, S.N. to	be over 210						
titre, 30 to 35° C			60 lb.	25	lb.	20	lb.
Wyo Tol No 710 (Callaide	I Dontonita		00 10	0	11.	10	10.
Wyo-Jel No. 719 (Colloida	1 Dentonite),	04 17	00 11	0.0	77	00	77
200 mesh		24 lb.	33 lb.	30	lb.	20	lb.
Tri-Sodium Phosphate, tech	1. grade pow-						
dered		1 lb.	7 lb.	. 5	lb.		
Perfume							
		0.2 lb					
41.							

0.1 lb.

0.7 lb.

0.1 lb.

Girella Camphory Sassafras Oil

^{*} c.p.s. = Chemically Pure Soap. † S.N. = Saponification Number. ‡ Titre = Melting Point of Fats.

The ingredients are weighed into a clean and dry mixer and intensely mixed for 15 to 20 minutes. The perfume should be sprayed or sprinkled over the powdered soap or soap-powder to avoid caking. As none of the ingredients are hygroscopic it is not necessary to pack

the finished product air tight.

For starting production, a clean openhead steel drum rolled and shaken on the floor is satisfactory for mixing, providing some wooden weights are laid inside to assure agitation. However, for big scale production, use one big horizontal mixer, 2000 lb. capacity, cylinder driven from both end countershafts and equipped with a double action agitator which moves toward the 6"x8" outlet in the middle and which is driven by a 15 h.p. motor. A slip ring motor, or a compensator allows this mixer to be started with a full load, thus avoiding accidents and dusting.

The most ideal process to make powdered hand toilet soaps is by making them wet-processed, and if other soaps are also manufactured, it is easy and much more preferable to do so. In the case of Formula 1, the Wyo-Jel is crutched into the hot molten soap stock before cooling and drying and the perfume is added immediately before grinding down of the dried soap flakes. In case of Nos. 2 and 3, paste soap, regular soap-powder is hot mixed with all the ingredients added at once to a bakery-type dough mixer. In case of hot processing much more Wyo-Jel can be used and the final structure will be more uniform and much harder to duplicate.

Liquid Soaps (French) Formula No. 1

Olive Oil Soap:

(Caustic Potash (Solid) Water

minimum possible for solution

b. Olive Oil	182 kg.
Palm Oil	362 kg.
Coconut Oil	362 kg.

Heat to 49° C., add to a.

c. Alcohol 170 l.

Boil the whole under reflux (82° C.). When saponified, cool, and add

. Water 5.6 l.

No. 2

Coconut Oil Soap

a. Soda Ash	1 kg.
Water	10 1.
b. Wood Ashes	15 kg.
Water	10 l.

Extract through a tin can with holes, pouring through water 3 to 5 times.

c. Caustic Soda

1. Boil 10 to 15 min.:

1 part by volume b. 4 parts by volume

6 parts by volume c. Add Coconut Oil 10 parts by volume

during the boiling in small parts, stir slowly. Then diminish heat, stir continuously, take off, stir, then pour into wooden forms.

2. Or: Boil 10-15 minutes:

Tallow

ъ. 4 parts by volume 6 parts by volume Sodium Sulphate

(10%)1 part by volume Salt 1/2 part by volume

Add: Coconut Oil 9 parts by volume and after:

Method as in No. 1. Gentle boiling, thorough stirring, dry.

1 part by volume

No. 3

Liquid Coconut Oil Soap

a. Water Caustic Potash (Solid) kg. Add a to

b. Coconut Oil (49° C.)

kg. c. Alcohol 2.5 I.

Warm the whole to 82° C. under reflux as in 1. Let cool 24 hours, then add:

d. Water 80 1. Sugar very little Potassium Chloride Glycerin optional

No. 4

Liquid Glycerin Soap

Soft Soap, Good	35 g.
Glycerin	21 g.
Water	7 g.
Alcohol	14 g.
Talc or Pumice	5 g.

Let stand for several days; take care to eliminate excessive alkali by adding oleic acid. Filter.

Transparent	Glycerin	Soaps
	Form	ula.

	No. 1	No. 2	No. 3
Coconut Oil, Cochin Tallow Castor Oil	20 18 12	26 24 10	30 kg. 20 kg. 15 kg.
Caustic Potasl 40° Bé. 36° Bé. 39° Bé. Glycerin	25 — — 10	$\frac{-}{32}$ $\frac{-}{13}$	— kg. — kg. 35 kg. 10 kg.
Sugar Water (60° C ''Fillers''	10 .) 15 —	40 30 30	42 kg. 38 kg. 35 kg.

To this soap-base add distilled water in small portions to about 15 (kg.), and to the resulting clear, but very soft, soap add a hardening solution (of 15° Bé.), made up of:

Potassium Carbonate	1 kg.
Sal Soda	1 kg.
Salt	1 kg.

Add water to get 15° Bé. Warm to

Add enough to get samples of sufficiently hard soap. Let stand covered for an hour, and test result.

Should not be of too high viscosity when spread on a glass-sheet. If too viscous or too foamy add water.

Add perfume at 50° C., sift in dye,

stir and pour into molds.

Transparent Soap (Without	Glycerin)	
Tallow, Cochin	24 kg.	
Coconut Oil	24 kg.	
Castor Oil	16 kg.	
Heat to 50-60° C.		
Add in thin jet:		
Caustic Soda (39° Bé.)	33 kg.	
Stir until soap swims on	top, ther	a
cover. Stir slowly over water	bath. Add	1
Alcohol	1-2 kg.	
then	-	
Water (60° C.)	22 kg. 20 kg.	
Sugar	20 kg.	

Again Alcohol 18-19 kg. Cover. Keep at 75° C. for an hour.

Soap should be dark and clear; foam light. Soap should remain "knifethick' on a glass-sheet.

If opaque, try (before in test-tube) to add slowly hot water, or caustic soda (20° Bé.).

At 50-60° C. add perfume and the last 3-4 kg. of above alcohol.

Rose Soap

a.	White Tallow Soap 1	0,000	g.
	Cinnabar, Moistened	68-08	ğ.
b.	Rose Essence	25	g.
	Geranium Essence	60	g.
	Clove Essence	15	g.
	Chinese Cinnamon Essence	10	g.

Polm Soan

	rain soap		
a.	Pure Palm Soap	5000	g.
	Half Palm Soap	5000	g.
b.	Bergamot Essence	60	g.
	Chinese Cinnamon Essence	25	g.
	Clove Essence	15	
	Essence of Fine Lavender	20	

Althaea (Marshmallow) Soap

5000 g.

5000 g.

3 g.

10 g.

a. White Tallow Soap

Pure Palm Soap

b. Yellow Ochre	30	g.
Paris Red	30	g.
c. Essence of Fine Lavende	r 15	g.
Essence of Pressed Lemon Peel Essence of Neroli	16	g.
Petitgrain Essence of Verbena	16 10	g. g.

Bouquet Soap

Essence of English Mint

α.	Soap, White Tallow Brown Ochre	10,000	g.
ъ.	Essence of Bergamot	80	g.
	Essence of Cloves	15	
	Essence of Neroli	15	g.
	Essence of Sassairas	10	g.
	Essence of Thyme	10	$\mathbf{g}.$
or	also:		
ъ.	Essence of Fine Lavende	r 20	g.
	Essence of English Mint	20	ğ.
	Essence of Pressed		
	Lemon Peel	25	g.
	Essence of Sage	20	g.

The following Soaps using Lauryl Sulphonates are covered by German Patents.

Essence of Thyme

I. True Lemon Soap

Citric Acid Sodium Citrate	5 g. 1 g.
Lanolin-Vaseline Oil (2:1	8-
1:1)	, 5 g.
Vegetable Lecithin	2 g.
Glycerin	2 g.
Lauryl Sulphonate	85 g.

II. I	iauid I	Car Soar)	1		v. 0	Chlorthy	mol i	Soap	
Wood Tar (1	•	^	3	g.	Chlor	othymo	. •			g.
Glycerin			5		Aceti	c Acid	, Conce	atrat	ed 0.5	g.
Triethanolam		uryl			Alcol				3.5	g.
Sulphonate		_	92	g.		hanolai lphonat	mine La e	uryi	95	g.
	. Alum	~	_	1		VI	. Chlori	ne S	oap	
Aluminum S			5	g.	Chlor				1-2	g.
Lorol Sulpha ethanolami							affin Oi	(1	:1) 5	g.
phonate	110 1300	1,71 Nu1	95	g.	Glyce		ryl Suln	hona	te 90	g. or
•	w 11	~		•	1300110	mi-rau	iyi buin	11()11(1	_	8.
	. Iodin	-	_	1	Q.o.	e 1	Damarin	Cl.	- manal Chi.	
Iodine-Alcoho	ol Solut	tion	5					g Do	earred Skir	
Glycerin Triethanolam	ine La	arvl	10	g.		d Para	oap, Po	wdar	70 e ed 70 g	
Sulphonate		ur, r	85	g.					up to 10	r.
								, -		,
			POW		ORMULA	E				
				Mag- nesium	Mag-					
	Rice	771	Colloidal	Car-	nesium	Zinc	Cold		Other	
Face Powder:	Staren	rateum	Kaolin	bonate	Stearate	Oxide	Cream		Additions	
	600	200 300	• • •	100	40	60				
	450 500	300		$\frac{50}{25}$		$\frac{220}{150}$		70 7	ritanium Die	oxide
D. J. D	500	300	100	250	5	• • •	• • • •			
Body Powder:		900				90		10	Salicylic Aci	d
	70	800 850		70	20	10		100	Boric Acid	-
	80	490	300		100		• • •	60 .	Boric Acid	
Infant Powder		1000								
Foot Powder:		1000		• • • •		• • •	ก	1.	Lanolin	
		850				100		(1/) Collegio A	.:.
	• • • •		• • • •	• • •	• • •	100	• • •	1 20) Salicylic A) Boric Acid	cia
		800 750				$\frac{200}{200}$		350	Boric Acid Kieselguhr	
		600						1 10	Thymol	
									Formaldeh	yde
		owders			***	.1 6	No.		_	
Phenol	ormula	NO. 1	7	~		uth Su : Acid	bgallate	,	5	g.
Camphor			3	g.	110011	Acid	No.	7	15	g.
Exsiccated A	lum		96	g.	Bism	nth Su	bnitrate		20	O*
	No.	9			Stare		***************************************		10	g.
Colinglia Ani		4		-	Purit	fied Ta	lc		70	g.
Salicylic Aci Boric Acid	CI .		5	g. g.			No.	8		
Starch			16	g.	Merc	uric Cl	ıloride		0.06	or.
Purified Tal	e			g.		ım Sal			26	g.
	No.	3			Prep	ared Cl	nalk		4	g.
Salicylic Aci			10							
Bismuth Sul		,	15	g.	T	hiosulp	hate D	istin	g Powder	
Zinc Stearat	е		10	g.	Sodi	ım Thi	osulpha	te	6	g.
0.71	No.	4		. 1	Borio	Acid	-		24	g.
Salicylic Aci	d		2	g.			owder	(pro	phylactic)	for
Tannoform Talcum			13 15	g.	ringwo	rm.				
			0	ρ,						

No. 5

2 g. 5 g. 33 g. 60 g. Foaming Bath Powder

40 g. 50 g. 10 g.

Sodium Acid Carbonate Starch, Wheat Sodium Carbonate

Salicylic Acid Tannic Acid Orris Root Alum

96	THE CHEMICAL	L FORMULARY	
Tartaric Acid	30 g.	Clove Oil	5 ec.
Kaolin, Colloidal	20 g.	Fennel Oil	5 cc.
Soap Powder, Concent	rated 45 g.	Ceylon Cinnamon Oil	1 cc.
Saponin	5 g.	Lemon Oil	1 cc.
	-	Demon On	1 00.
Keep completely dry a	ind sealed from		
air to avoid decompositi	on. 1-2% per-	Oxygen Tooth Pa	ste
fume (Lavender, Pine	Needle, Eau de	Calcium Carbonate, Preci	pi-
Cologne, Fancy), is added	1.	tated, Medium Density	40 g.
		Glycerin, C.P.	30 g.
Mentholated To	alcum	Hard Fat Soap Powder	7 g.
Menthol	0.25 g.		1 soft paste
Alcohol	5 cc.	To 100 parts of this pas	
Talcum	50 g.		
	9 1	Sodium Perborate Perfume	10-15 g.
Dust freely on itching	part.	Feriume	1 g.
"Prickly Heat"	Powder	Tale Tooth Pas	ste
Starch	12½ lb.	Purified Tale	42 lb.
Talc	7 lb.	Magnesium Carbonate	8 lb.
Zinc Stearate	½ lb.	Phenol	½ lb.
Camphor	2 oz.	m	
Zinc Oxide	5 lb.	Oil of Orange Oil of Lemon Oil of Anise	214 dram
Menthol	1 oz.	Oil of Lemon	5 0%
and out of the control of the contro	2 021	Oil of Anise	1 dram
		Oil of Peppermint	6 oz.
Tooth Pas	te	Menthol	5 oz.
Soap Powder	2500 g.	Glycerin	6 gal.
Calcium Carbonate	500 g.	diyeenii	o gan
Lactose	150 g.		
Glycerin (28° Bé.)	2000 g.	Salt Tooth Pas	te
water	400 g.	U. S. Patent 1,96	8 858
Peppermint Oil	100 g.	•	•
Alcohol	100 g.	Glycerin, C.P.	$37\frac{1}{2}$ lb.
Carmine	10-20 g.	Glycerin, C.P. Neutral Soap Gum Tragacanth	1½ lb.
	_ ' '	Gum Tragacanth	1½ lb.
Tooth Paste with Low (Hyeerin Content	Magnesium Carbonate	
	-	Magnesium Carbonate (Finely Divided) Calcium Carbonate	13 lb.
Calcium Carbonate, Pr	ecipi-	Calcium Carbonate	
tated, Medium Dens		(Finely Divided)	$51\frac{1}{2}$ lb.
White Clay (Bolus Al	ba) 5 g.	Milk of Magnesia (Mag	• <u>.</u>
Soap Powder (85–88%)),	nesium Hydroxide)	31 lb.
Pale, no Odor or Tas		Distilled Water	24 pt.
Water	20 g.	Saccharin Powder	282 gr.
Glycerin, C.P.	20 g.	Salt (Finely Divided)	108 lb.
Tooth Paste (Witho	ut Glycerin)	Flavor	
White Clay (Bolus Calcium Carbonate,	Alha) 30 c	Menthol Crystals	$2\frac{2}{3}$ oz.
Calcium Carbonata	Alba) 50 g.	Oil of Peppermint, U.S.	P. 8 oz.
a. Precinitated	15 m	Oil of Anise, U.S.P. Methyl Salicylate *Flavor Compound	2/3 oz. 2/3 oz.
a. Precipitated Soap Powder (as A b. Tragacanth Paste (hova) 1 g.	Methyl Salicylate	2/3 oz.
h Tracacanth Paste	10%) until nagty	*Flavor Compound	,,,
o. magacantin raste (170) until pasty	No. 04595	12 oz.
×		*Flavor Compound No. 045	95 is comprised
Tooth Pas		as follows:	•
Calcium Carbonate, P.	re-	Twice Rectified Oil of	
cipitated	50 g.	Peppermint	274 oz
White Bolus	10 g.	Oil of Eucalyptol Oil of Wintergreen	90 oz. 45 oz.
Glycerin (sp. gr. 1.24,	30° Bé.) 20 g.	Rectified Aniseed Oil	22 1/2 oz.
Water	18 g.	Safrol	22 ½ oz.
Tragacanth	1 g.	The glycerin, water, son	p, gum traga-
Perfume (as below)	T	canth, milk of magnesia,	
Peppermint Oil	50 ec.	are mixed with a rapid m	
Peppermint Oil Menthol Anise Oil	50 ec. 5 ec.	Then flavor is added, w	

15 minutes of mixing the product is transferred to a small mixer, the salt is added, the mixer is run for five minutes more, then the magnesium carbonate is added, followed by another five minutes' run, after which the calcium carbonate is fed to the pasty mass, and, after this has been taken up, the batch is run for 20 minutes more.

The finished mass is allowed to stand for 12 hours, and, after stirring slowly for 10 minutes before filling, the mass is filled into ordinary collapsible tubes.

Denture	(Artificial	Teeth)	Clea	ner
	of Starch		36	g.
Diglycol			1	g.
Sugar Sy			2.25	g.
Magnesia	ım Carbona	te	1.13	g.
Gum Tra	igacanth			7 g.
	ted Chalk		41	g.
Sodium :	Bicarbonate		6	g.
Water			10.5	g.
Flavor			to s	uit

Denture (Artificial	Teeth)	Adherent
Gum Karaya		80 g.
Gum Arabic		20 g.

Dental Impression Material British Patent 399,842

	Copal	26	g.
	Stearic Acid	21	g.
	Shellac	5	g.
iı	Melt together and then add ag and stirring:	while	heat

Talc			48	g.
Iron	Oxide,	Red	48 1 <u>/</u> 4	g.

Temporary Dental Filling

Zinc Oxide	85	g.
Rosin, Powdered	15	g.
Oil of Cloves	60	g.
Canada Balsam	-35	g.
Peru Balsam	5	g.

Dental Canal Cement

Thymol	1	g.
Rosin	9	g.
Chloroform	150	g.

Dental Pulp Capping

Make a paste of zinc oxide and eugenol.

Dental Pulp Devitalizer

Make a paste of arsenic trioxide and eugenol

Antiseptic Mouth Wash ("Listerine", Type)

Boric Acid	50	g.
Benzoic Acid	1	g.
Thymol	1	g.
Eucalyptol	0.125	cc.
Oil of Peppermint	0.5	cc.
Oil of Wintergreen	0.25	cc.
Oil of Thyme	0.1	cc.
Grain Alcohol	250	cc.
Water to make up to		cc.
Caramel	to co	olor

The boric acid is dissolved in the water or about 700 cc. of same. All the other products are dissolved in the alcohol and the two solutions mixed and colored to a very pale straw. The above product must be labeled 25% grain alcohol.

Mouth Wash Tablets

Peppermint Oil	30	ec.
	100	
Sodium Benzoate	500	

Mouth Rinse

Salt	30	g.
Sugar	20	g.
Oil of Cinnamon	1/4	ec.
Oil of Cloves	1/2	cc.
Oil of Peppermint		ec.

Gingivitis Mouth Wash

Boric Acid	4 g.
Potassium Chlorate	8 g.
Peppermint Water	350 ec.

Breath Deodorant

Dissolve one 4.6 grain tablet chloramine in 1 oz. water. Brush teeth and tongue, and rinse out mouth with this solution, while fresh.

Immediately and permanently rids breath of even such odors as those of garlic and onions.

Depilatory

German Patent 601,078

Barium Sulphide	100	oz.
Starch	- 60	oz.
Magnesium Silicate	30	oz.
Pyrogallol	10	oz.

Make into a paste with water before using.

Odorless Depilatory

Perhydrol	3.5 - 5	ø.
Polychol (or Polyglycol)	5	
Lanolin Anhydrous	20	g.
Rub together till uniform.		Θ.

Adhesive Depilatory U. S. Patent 2,013,928 Rosin 90 g. Cottonseed Oil 10 g. Warm together and stir until uniform.	Sunburn-Protecting Oil Quinine Oleate, C.P. 3-5 g. Paraffin Oil 27 cc. Fatty Oil 70-68 cc. Dye (Oil-Soluble Red)
Sun Burn—Protectors Liquid a. Triethanolamine 40 g. Trihydroxyethylamine	Sunburn-Protecting Oil Quinine Ricinoleate 3-5 g. Olive Oil 97-95 cc.
Stearate 40 g. Melt on water bath, make emulsion in Water (60° C.) 620-630 g. b. Paraffin Oil 100 g. Peanut Oil 150 g. Oleic Acid 30 g. Warm up on water bath to 40° C. Methyl-p-Hydroxy Ben-	Sunburn (Suntan) Oil Mix Vaseline Oil 75 g. Sesame or Peanut Oil, Pale 23 g. Thymol 0.5 g. Lanolin, Anhydrous 1.5 g. Perfume 1-2 g. Made up of:
zoate 1 g. Pour b into a, perfume with c. Perfume Oil to suit Stir until cold.	Pine Oil 3 cc. Lavender Oil 1 cc. Rosemary Oil 1 cc. Laurel Oil 3-5 cc.
Over	Suntan Oil
Cream White Wax 60 g. Cocoa Butter 30 g. Lanolin, Anhydrous 40 g. Peanut Oil 300 g. Spermaceti 20 g. Moldex or Other Preservative 1 g. Perfume 5-10 g.	Paraffin Oil 20 cc. Fatty Oils, Free from Acid, Preserved 80 cc. Etheric Oils (Bergamot, Eau de Cologne [free from Methylanthranilic Ester] or Pine Needle Oil) 1 cc. Dye with Chlorophyll, Oil-soluble.
Preventatives against Sunburn a. Gum Tragacanth (Powder) 15 g. Glycerin 50 g. Grind in mortar.	Preparations to Protect Feet Agains Hurting and Inflammation Foot Creams
b. Quinine Acid Sulphate Citric Acid Water Alcohol (95%) with Perfume c. Glycerin Grind a, then add the b solution, and	Formula No. 1 Potash Soap 50 g. Yellow Vaseline 15 g. Water 29 g. Zinc Oxide 6 g. Caustic Soda 11 drops No. 2
Sunburn Protecting Cream a. Quinine Hydrochloride 4 g.	Potash Soap 52 g. Vaseline 15 g. Water 27 g. Zinc Oxide 6 g.
a. Quinine Hydrochloride 4 g. Alcohol (95%) 12 g. b. Citric Acid 0.8 g. Water 10 g. c. Tragacanth Powder 3.5 g. Glycerin 10 g. Water 42.5 g.	No. 3 Soap 35 g. Vaseline 15 g. Water 45 g. Zinc Oxide 5 g. Lavender Oil to suit
Mix solutions a and b and then work into solution c. Perfume Composition, with Fresh Perfume Odor 9 drops	No. 4 Lamb Tallow 100 g. Pig Fat 100 g. Creosote 1 g. Juniper Oil 10 g.

No. 5	
Wool Fat	20 g.
Vaseline	10 g.
Formalin	10 g.
No. 6	
Glyceryl Monostearate	20 g.
Glycerin	5 g.
Paraffin Oil	5 g.
Formaldehyde Solution	15 cc.
Water	55 cc.
Melt up to 60° C. Stir until	cold.

Peeling Paste for Corns or Hard Skin (Not to be put on normal skin, as it is irritating).

Formula No. 1

Lard Salicylic Acid, U.S.P.	50 g. 50 g.
No. 2	
Salicylic Acid, C.P.	30 g. 70 g.
Vaseline, White	70 g.
No. 3	

Mild-acting paste (stir warm):

mind-acting paste (stir	war	uı):
Pine Resin, Pure	8	g.) 353
Wax, Yellow	30	$\left\{ \begin{array}{l} g. \\ g. \end{array} \right\}$ Melt
" Larch Turpentine	12	g.
^l Vaseline, Ÿellow	16	g.
b. Salicylic Acid	8	g.
Anaesthesin	3	g.
Peanut Oil	14.5	g.

Mix warm, stir until clear solution; cool stirring; when thickening starts, add Methyl Salicylate 0.5 g. Peru Balsam 8 g. Stir until cold

Athlete's Foot Ointment

	oz.
Ammoniated Mercury 4	oz.
	oz.
Oil of Eucalyptus 12	oz.
	oz.
Mix and make into an ointment.	

Athlete's Foot Powder

Sodium Thiosulphate	20	oz.
Boric Acid	50	oz.
Purified Talc (Sterilized)	30	oz.

Triturate thoroughly. This may be used as a prophylactic powder applied to the feet and dusted in the shoes.

Athlete's Foot Treatment

Immerse feet two or three times a day in a warm saturated aqueous solution of furfural. Always have a little free furfural floating around to make sure of an excess. Continue treatment until all signs of the disease disappear. Then treat feet once a day for several weeks to prevent recurrence. Shoes and socks should also be treated with this solution to disinfect them.

"Athlete's Foot" Remedy

TECHNOLO D T OOC	account of
Gentian Violet	1 part
Alcohol	100 parts
Water	100 parts
O	•

Stir until dissolved.

Bunion Remover

Salicyl	ic Ac	id			6	g.
Lanoli	n					g.
Soak f	oot in	hot	water.	out	off	thial

soak foot in hot water; cut off thick skin and apply twice a day.

Pilocarpine Eye Drops

Pilocarpine	Nitrate	•	0.1	σ.
Boric Âcid			0.2	
Distilled Wa	ater	to make	e 10	cc.
Label: Drop	into eve	from o	ne to	five

times daily (in chronic glaucoma).

Pilocarpine Eye Salve

Pilocarpine Nitrate	0.2 g.
Distilled Water	1 cc.
Hydrous Wool Fat	2 g.
White Petrolatum	7 g.

Mix with careful trituration and dispense in collapsible tube with eve tip.

Label: Apply to affected eye at bedtime (in chronic glaucoma). If collapsible eye ointment tube is not available, a glass rod may be used to apply salve to lower lid, which is then permitted to close. Gentle massage of lids helps to distribute ointment over the conjunctiva.

Eye Ointment

	-		
Silver Nitrate		0.5	g.
Distilled Water		1	g.
Cocoa Butter		15	g.
Liquid Paraffin	7	equal pa	rts
Soft Paraffin	1	to 100	g.

Cetyl Alcohol U. S. Patent 2,021,926

Formula No. 1

241 parts of spermaceti are melted and heated to 200° C. 42 parts of powdered potassium hydroxide are now added with agitation in half an hour, during which time the temperature is allowed to rise to 240° C. It is held at this temperature for half an hour when superheated steam

solved.

is passed in. There distils over with the steam a colorless oil which sets on cooling to a crystalline waxy solid which is entirely free from fatty acid and from unsaponified spermaceti. The yield is approximately 100 parts by weight, the proportion of water to oil in the distillate being approximately 10:1.

No. 2

241 parts of spermaceti are treated as in Example 1 with a mixture of 21 parts powdered potassium hydroxide and 15 parts of powdered sodium hydroxide. After reaction, the molten mixture of soaps and fatty alcohol is subjected to superheated steam distillation at about 250° C., eventually at 280° C. until no more oil distils. The yield is approximately 100 parts of the pure alcohol from spermaceti, the ratio of water to oil in the distillate being approximately 10:1.

No. 3

268 parts of sperm oil are treated as in Example 1 with a mixture of 21 parts of caustic potash and 15 parts of caustic soda. After reaction the mass is subjected to superheated steam distillation until no more oil distils. The yield is 90 parts of a semi-solid alcohol, free from unsaponified wax or free fatty acid. The ratio of water to oil in the distillate is approximately 4:1.

Arthritis Ointment

Ichthy Lanol			20 30	g. g.
				_

Rub together until uniform; apply freely to joint and apply bandage.

Frostbite Ointment

Ichthyol	3 g.
Lanolin	4 g.
Camphor	2 g.
Petrolatum	60 g.

Warm and stir until dissolved. Rub into skin and bandage.

Analgesic Balm

Menthol	5 oz.
Methyl Salicylate	10 oz.
Hydrous Wool Fat	75 oz.
White Petrolatum	10 oz.

Burn Ointment

Tannic Acid	2 g.
Ichthyol	33 g.
Lanolin	62 g.

Carbuncle Ointment

Ichthyol	25 g.
Lanolin	35 g.
Zinc Oxide Ointment	90 g
Apply thickly daily.	

Chapped Skin Ointment

Phenyl Salicylate		8 g.
Menthol		4 g.
Olive Oil		40 cc.
Lanolin		125 g.
Warm together a	nd mix	until dis-

Glycerin-Sulphur-Kaolin-Acne Paste Kaolin 10 g. Sulphur, Colloidal 7.5 g.

Glycerin (24%) to pasty consistency

Boil Ointment

Ichthyol	15	g.
Lanolin	68	

Apply thickly on gauze and hold in place with adhesive.

Ringworm Ointments Sulphur Ointment

ourbreak ourrestone		
Precipitated Sulphur	1.5	g.
Petrolatum	30	g.

Rub in gently once or twice daily. Strength may gradually be increased up to 20 per cent.

Compound Benzoic Acid Ointment

Salicylic Acid	1 g.
Benzoic Acid	2 g.
Ointment of Rose Water	30 g.
Apply locally twice dails	v. Strengt

Chrysarohin Ointment

may be doubled, if necessary.

CHIJBAIODH	Omment	'		
Chrysarobin Petrolatum		$\begin{array}{c} 1.5 \\ 30 \end{array}$	g.	

Apply with care against getting it in the eyes.

Salicylic Acid Pigment

Salicylic Acid	1.5	g.
Chloroform		cc.

Paint on affected area twice daily until desquamation occurs.

Pyrethrum Ointment

Pyre	thrun	Extrac	t		27	g.
Abs	orption	n Base	(Parachol	()	73	g.
Mix	until	smooth.	Useful	in	tre	atin

scabies and other insect infestations.

Ulcer Salve

Ethyl Aminobenzoate	3 g.
Paraffin	10 g.
Petrolatum	20 g.
Spread on cause and apply to	ulcer.

Protecting Skin Against Mustard Gas

Glycerin impregnated coarse fibered clothing is recommended. This protection lasts for at least two hours' exposure to this gas.

A. B. C. Liniment

Tincture of Aconite	30	cc.
Fluidextract of Belladonna	30	cc.
Chloroform	30	cc.
Soap Liniment to mak	e 240	cc.
Analgesic liniment. For e	xterna	l use

only.

Glycerin-Sulphur Liniment

53	
Potassium Carbonate	20 g.
Glycerin	20 g.
Sulphur, Precipitated	20 g.
(Grind together)	•
Alcohol (68%)	20 g.
Ether	20 g.

"Penetrating" Liniment

	T 01100110011119			
Oil of	Turpentine		1	gal.
Oil of	Sassafras		1	lb.
Oil of	Cajaput		1	lb.
Chloro				gal.
Oil of	Camphor		1/4	gal.
Oleores	sin Capsicum		5	oz.
Coal O	il -		3	gal.
		-		-

Rheumatism Liniment

Camphor	1 lb.
Chloroform	32 fl. oz.
Alcohol	80 fl. oz.
Methyl Salicylate	16 fl. oz.

Dissolve camphor in the mixture of the other ingredients. Excellent for sore or aching muscles. Should be applied at night by rubbing in.

Back Rub Ointment

Zinc Stearate	5	g.
Tincture of Benzoin	5	ğ.
Scarlet Red Ointment		g.
Hydrous Wool Fat	30	g.
Liniment of Camphor	180	cc.
Mutton Tallow	500	g.

Non-Staining (Non-Leaking) Mineral Oil Laxative

White Soft Paraffin Wax	2 oz.
Mineral Oil, U.S.P.	6 oz.
Warm together and stir until	uniform

Castor Oil Candy Laxative U. S. Patent 1,991,139

Predetermined quantities of broken chocolate and castor oil are heated in separate containers or kettles before mixing. The chocolate is heated to approximately 115° F., while being thoroughly stirred or agitated, and is then permitted to cool to approximately 85° F., which temperature is finally slowly increased to between 88 and 90° F.

After the chocolate melting operation has been commenced, or simultaneously with this operation, an amount of castor oil approximately that of the melted chocolate, is slowly heated to between 85 and 90° F., preferably between 88 and 90° F. The heating of the castor oil and chocolate is so timed that the temperature of the one will coincide with that of the other. The best mixing temperature is between 88 and 90° F., it being essential that the temperature of each ingredient be kept exactly the same.

Mixing of the melted chocolate and heated castor oil is effected at this stage by drawing off the two ingredients from their separate kettles into a mixer, where they are thoroughly beaten and blended, after which the temperature is lowered to between 75 and 80° F. At this point, the mixture is cast into centers or chocolate shells which are subsequently capped with chocolate and run into a cold box for final cooling.

Agar Mineral Oil Emulsion

Mineral Oil	~	$18\frac{3}{4}$	gal.
Emulsone B Tragacant		834	lb.
Powdered A		1	lb.
Citric Acid		2	OZ.

Some sodium benzoate and aseptoform as preservative, and a small amount of vanillin and saccharin for flavoring purposes.

Emulsion of Liquid Petrolatum

	- COLOTONOUS	
Liquid Petrolatum	500	cc.
Acacia, in Very Fine		
Powder	125	g.
Syrup	100	ec.
Vanillin	0.035	g.
Alcohol	60	cc.
Distilled Water, a suffic	cient	
quantity to make	1000	cc.

Mix the liquid petrolatum with the powdered acacia in a dry mortar, add 250 cc. of distilled water all at once and emulsify the mixture. Then add, in divided portions and triturating after each addition, a mixture of the syrup, 50 cc.

of distilled water and the vanillin, dissolved in the alcohol. Finally add sufficient distilled water to make the product measure 1000 cc.

Note: In preparing Emulsion of Liquid Petrolatum other methods of emulsification may be used and the quantity of acacia may be reduced or it may be replaced by agar, gelatin, tragacanth or mixtures of any of these emulsifying agents, provided the resulting emulsion is similar in viscosity and appearance to the emulsion made by the formula suggested above.

Antipyrine Suppositories

Antipyrine	3 g.
Extract of Belladonna	0.1 cc.
Cacao Butter	20 g.

Mix and divide into ten suppositories. Label: One every two to four hours as required.

Psoriasis Treatment

Formula No. 1

Salicylic Acid			10	g.
Oil of Cade			25	cc.
Soft Soap			25	g.
Alcohol	to	make	100	čc.

Paint over patches, permit to dry, and wash off excess in bath.

No. 2

Salicylic Acid	10	g.
Chrysarobin	20	čc.
Oil of Cade	20	cc.
Soft Soap	25	g.
Petrolatum	25	g.
T 7 7 4 7		_

Label: Apply to patches.

Acidosis Preventative

To a teaspoonful of sodium bicarbonate in a deep bowl, add the juice from one lemon. Stir until effervescence is completed, and add a glass of cold water, and drink. Best results are obtained by taking this drink upon rising in the morning, at least one-half hour before breakfast.

Cold and Grippe "Remedy"

The following has been used with splendid success by members of a technical manufacturing organization:

a. Acetic Acid (36%),

U.S.P. 1/3 fl. oz. Water to make 1 fl. oz.

b. Ammonium Carbonate, U.S.P. 48 gr. Water to make 1 fl. oz. c. Sodium Bicarbonate 2 d.
Potassium Citrate 2 d.
Aromatic Spirits of
Ammonia 1 fl. oz.
Water 1 fl. oz.

Mix a and b; after effervescence stops add c.

Take one teaspoonful every 2 hours.

Hay Fever Remedies Formula No. 1

Ephedrine (Dried) Petrolatum, Liquid	$\begin{array}{cc} 0.1 & \mathrm{g.} \\ 10 & \mathrm{cc.} \end{array}$
Use as nasal spray.	

No. 2

Ephedrine Sulphate 1 g. Calcium Lactate 4 g. Place in No. XXX capsules; use one

3 or 4 times daily.

Sea-Sickness Remedy

Antipyrin	4 g.
Sodium Bromide	8 g.
Sugar	2 g.

Use once every three hours.

Appetite Stimulant

Appente Simulant		
Tincture of Capsicum	2	cc.
Tincture of Nux Vomica	16	cc.
Tincture of Gentian		
Compound	72	cc.

Dose: Three teaspoonfuls daily.

Bronchitis Inhalant

Menthol		1/2	g.
Chloroform		4	cc.
Tincture of	Benzoin	120	cc.

Inhale twice daily, using one teaspoonful to pint of boiling water.

Menthol Inhalator

Eucalyptus Oil	4	cc.
Menthol	2	g.
Paraffin Oil	94	čc.

Laryngitis Spray

Thymol		0.15	g.
Menthol		1.2	g.
Eucalyptus Oil		3	g.
Petrolatum, Liquid	3	00	ec.

Tonsilitis Gargle

Potassin	um Chlo	rate	- 8	g.
		Chloride	12	cc.
Glycerin	1		60	cc.
Water			240	cc.

Stomach Gas Relief Calomel Bicarbonate of Soda Lactose	3 g. 5 g. 4 g.
Periodic Pain Allevia Formula No. 1	tor
Amidopyrine Alcohol Simple Syrup Flavor No. 2	20 oz. 40 oz. 138 oz. to suit
Starch Amidopyrine Acetyl Salicylic Acid	90 oz. 90 oz. 25 oz.
Camphor Tablets (Pharmac	ceutical) 5 g.

Pack tight, to prevent volatilizing. Moth Protection Tablets

Peppermint Oil

Naphthalene Camphor Ceresin	225 75 50	g.
Melt together and then add		
Hexachlorethane	50 5	g. g.

Dip cardboards into the above while

Sterilizing Helmets and Gas Masks

The U.S. Government, in its specifications for sand blast helmets purchased by its various departments, requires that each article be capable of passing either one of the following sterilization tests:

(a) Immersion for ten minutes in a solution of formaldehyde made by placing one part of 40% solution of formaldehyde in nine parts of water, or

(b) Subjection to sterilization by a moist atmosphere of antiseptic gas, preferably formaldehyde, for a period of ten minutes, at room temperature.

It has been suggested that care should be taken to remove all the formaldehyde from the masks by washing with water before they are placed in use.

"Creolin" Disinfectant

Sulphonat	ed Caston	Oil	100	kg.
Caustic Sc	da (36°	Bé.)	51.2	
Heat abov	re at 80-	L00°C.,	then a	d
Rosin			104	
Mix with	heating	until	uniforn	n and
add	St. 19.57			
Tar Oils	200-3209	(C)	775	kø.

Mix until	dissolved	and	then	add
Water	to	make	1000) kg.

Disinfectant for Telephones

Solution 1

ROZGIJA Z	
Oil of Wintergreen	0.5 g.
Oil of Eucalyptus	0.25 g.
Denatured Alcohol	15 g.
Solution 2	

Formaldehyde	25	cc.
Water	225	cc.

Add solution 1 to solution 2 and dilute with water to 1000 cc.

Counter Irritant, Extra Strong

Menthol					2 g.
Volatile	Oil	\mathbf{of}	Mustard		cc.
LadaalA				= (٠

Apply a few drops to affected area. (Must not be used in the vicinity of the

Stainless Iodine Solution

Resublimed Iodine	4 g.
Potassium Iodide	10 g.
Hyposulphite of Soda	10 g.
Alcohol, Anhydrous	200 cc.

Non-Irrititating Iodine Antiseptic

Iodine	2 g.
Potassium Iodide	2.4 g.
Alcohol	55 g.
Water	45 g.

Tattoo Mark, Removing

First the skin is vigorously rubbed until the outer epidermis comes off; then a paste of quicklime, just slacked, to which pulverized phosphorus (two tablespoonfuls to a pint) is added and thoroughly mixed, is applied to the tattooed surface and held by a bandage, which is taken off two days later. The crust is left to dry and then fall off itself; in about fifteen days. A second application should be made; a third is rarely necessary. Thus treated, the tattooing disappears completely without the least scar.

Mechanics Hand Protective Coating II S Patent 2021 131

O. D. Latent 2	1,011,101
Water	1600 oz.
Sodium Stearate	288 oz.
Glycerin	1155 oz.
Sodium Silicate	906 oz.
Lemenone	1 oz.

Volatile Anæsthetics Formula No. 1

Methyl and Ethyl Chloride equal parts

No. 2			
Ethyl Chloride		60	cc.
Methyl Chloride		35	cc.
Ethyl Bromide		5	cc.

No. 3

Methyl Chloride Ethyl Chloride Chloroform	}	equal	parts
---	---	-------	-------

An anæsthetic for external use containing

Chloroform	1/2	fi.	dr.
Ether	$2\frac{1}{2}$		
Liquid Paraffin	$2\frac{1}{2}$	fl.	dr.
laws J web on limb+	annatha		ia *

is employed when light anæsthesia is required in painful wound dressings or for short operations.

Anæsthesia Chloroform Preservative Add 1% of absolute alcohol and keep in a cool place away from direct light.

X-Ray Contrast Media

1. Barium diet for stomach and intestinal examinations. Boil together

OD CERTOS CAROLINA INC. CALONIO		
Corn Starch	15 g.	
Sugar	15 g.	
Cocoa	20 g.	
Barium Sulphate	150 g.	
Water	500 cc.	

2. Barium diet for diagnosis of stenosis of the small intestine.

Bariu Thick	m Sulphate Gruel		80 g. 200 cc.
		• .	

3. Barium suppository for rectum examination.

Corn Starch		30 g.
Water	750	to 1000 cc.
Boil together and	add a	suspension of
Barium Sulphate		200 g.
Water		500 cc.

Cystographic Medium U. S. Patent 1,935,661

Five to 8 per cent aqueous solutions of sodium (or potassium) bismuth tartrate or citrate (1) serves as cystographic media opaque to X-rays; (1) should contain about 65 (70) per cent of bismuth.

Hormone Manufacture U. S. Patent 1,978,297

The ground whole testicles are preferably macerated from 12 to 48 hours with

the required amount of the solvent selected, the liquid is filtered off, the residue expressed and re-extracted with preferably the same solvent, this time (the glands having been freed from the water therein) using the exact concentration which recovers most of the hormone with the least undesired material, as, for example, 90% acetone, 70% propyl alcohol or about 75% ethyl or methyl alcohol, Extraction is continued until the residue is fully extracted. The extracts are combined, and the solvent distilled off at a low temperature and under reduced pressure. All traces of the solvent are removed, leaving the lipoid material containing the hormone, together with other substances emulsified in an aqueous solution.

The mixture resulting from the agitation of the emulsified aqueous solution of the lipoid material with one of the solvents named above, when the agitation has ceased, separates into two or three layers, dependent upon the solvent used. When three layers are formed, the upper or solvent layer contains the active lipoid with possible traces of cholesterol and phospholipins, and is free from protein, the middle layer contains most of the phospholipins and cholesterol present in the original extract, together with other organic material, and a portion of the solvent and water. The lower aqueous layer contains blood pigments, salts, etc. The one or two lower layers are preferably drawn off and the agitation with the hormone solvent repeated several times and finally the two or three layers are drawn off separately. In case chloroform is used the lower chloroform layer contains the active hormones.

The combined upper layers may then be washed with a 1 to 10% sodium carbonate solution to remove all traces of the fatty acids and phospholipins, washed with water to remove the sodium carbonate and the solvent distilled off. The residue then contains the testicular hormone in a high state of purity.

For example, in using amyl alcohol at this step of the process, the agitated mixture of the amyl alcohol and the aqueous solution containing the lipoid material separates into three layers, with the upper layer containing the active portion or hormone. The two lower layers are then drawn off, the agitation with amyl alcohol repeated and the upper layers resulting from several repetitions of this step combined, washed with a 1 to 10% sodium carbonate solution and then with water and the solvent distilled off leaving the hormone in a high state of purity.

Analgesic Chaulmoogra Oil	for Injection
Chaulmoogra Oil	80 cc.
Olive Oil	20 cc.
Benzyl Ephedrine Base	0.1 g.

Intravenous Colloidal Sulphur British Patent 433,833

Sodium Sulphide, Pure	23.5	g.
Water, Distilled and Deaerated	50	cc.
Dextrin Dissolved in	10	g.
Water, Distilled	400	cc.
Dilute to	1	l.

Add sulphur dioxide to a pH of 7.6 and dilute with distilled water to 10 mg. of sulphur per ec.

Hydrogen Peroxide Preservative

The addition of 20 g. phenacetin to 5 kg. hydrogen peroxide acts as a good preservative.

Preservatives for Hydrogen Peroxide

According to French chemists the best preservative for hydrogen peroxide solution is phenetidine lactate in the proportion of 0.5 g. per liter of solution. Less effective are glucose, gelatin (0.2 g. per liter); ethyl alcohol (16 g. per liter); and hippuric acid (0.2%).

Embalming Fluid—For Decolorizing Jaundice Cases

U. S. Patent 1,942,407

Benzoyl Peroxide	15	g.
Ethyl Alcohol (95%)	3	gal.
Formalin (40%)	4	pt.
Water	$1\frac{1}{2}$	gal.

Embalming Fluid Formula No. 1

T OTHER TION T		
Formalin (40%)	220	oz.
Glycerin	100	oz.
Borax	90	oz.
Sodium Chloride	10	oz.
Sodium Nitrate	10	oz.
Potassium Citrate	50	oz.
Methanol	40	oz.
Water	75	oz.
Benzaldehyde	6	oz.
Color with Erythrosine.		

No. 2

Borax			4	oz.
Phenol			5	oz.
Salicylic Act	id		5	OZ.
Formalin (4	.0%)		71	oz.
Glycerin			.31	oz.
Water	sufficient	to	make 1	gal.

Corpse Wound Filler

a. Yellow Beeswax	- 5	oz.
Paraffin	5	oz.
White Petrolatum	15	oz.
b. Soap Flakes	2	oz.
Water	5	oz.

Finishing Cream (Corpse)

0	\ <u>+</u> /	
Glycol Stearate	1	2 oz.
Glyceryl Tristearat	e	5 oz.
Rose Oil		2 oz.
Glycerin		3 oz.
Water	7	8 oz.
Titanium Dioxide		1 oz.

Animal Embalming Fluid

Use a water solution of either 5% furfural or 10% formaldehyde.

Air Purifier

Alcohol (95%)	2000	cc.
Formalin (40%)	400	cc.
Pine Needle Oil	190	cc.
Thyme Oil	10	cc.
For use dilute with water	1:50.	

Solid, Volatile Preparations to Perfume and Disinfect the Air

> Formula No. 1 Naphthalene, Pure

> > No. 2

Paradichlorobenzol, Pure

No. 3*

Naphthalene, Scales	70	g.
Camphor, Sublimed	10	g.
Paradichlorobenzol	20	ğ.

No. 4

Naphthalene		80	g.
Carbolic Acid	(Phenol)	20	

Heat the mixtures gently, very little beyond the melting point (color optionally with yellow, red, blue, oil-soluble dyestuffs) and pour into molds. Work in well ventilated rooms.

* 0.5% of Citral may be added.

Water Soluble Bactericide U. S. Patent 1,930,474

A 1:1 mixture (200 g.) of chlorothymol and olive oil is treated with sulphuric acid (60 g.) at 20° for 2 days, and then washed free from acid with saturated aq. sodium sulphate; the product is readily dispersed in water.

Protecting Tin Collapsible Tubes Against Corrosion

U. S. Patent 1,968,722

Collapsible tubes containing soap, shaving cream, toothpaste and other alkaline materials are protected against corrosion by addition of 0.1% sodium nitrite.

Pharmaceutical Charcoal Preparations Tablets

Formula No. 1	
Activated Carbon	200 g.
Gum Tragacanth	8 g.
Sugar	195 g.
Water	68 g.
No. 2	
Activated Carbon	100 g.
Sugar	5 g.
Albumen Solution	5 g.
Gum Arabic	3 g.
Tincture of Benzoin	1 g.

The above are useful in the treatment of dyspepsia.

Removing Creosote from Skin and Clothing

Wash with isopropyl alcohol to remove creosote and prevent further "burning" of skin.

Zinc Ointment

White Beeswax	60	g.
Spermaceti	60	ğ.
Oil of Sweet Almonds	300	g.
Digest 2 hours on water	bath.	_
Gum Benzoin, Siam	20	g.
Add while cooling.		_
Zinc Oxide	100	g.
Boric Acid	2	g.
Carmine enough	i to col	lor
Perfume with extract of ro	se leav	es.

Hiccough Remedy

Take one teaspoonful of tincture of castoreum and repeat in a half hour if needed.

Fingernail Cleaner

A fingernail stain remover consists of a saturated solution of tartaric acid in water.

EMULSIONS

GASOLINE EMULSIONS

Formula No	1		Oleic Acid	1	ec.	No. 17		
			Alcohol		cc.			
Triethanolamine	11/2		Gasoline	45		34% Wat		
Water	11/2	cc.	No. 9	10		Triethanolamine	175	
Oleic Acid	5	cc.	Triethanolamine	7	ec.	Water	260	
Butanol Gasoline	45	cc.	Water		cc.	Stearic Acid	1400	ec.
			Stearic Acid		cc.	Alcohol Gasoline	1400	
Dissolve trietha			Alcohol		cc.	Gasonne	31500	cc.
in water and add			Gasoline	45		No. 18		
ture of other in	gredie	nts	No. 10	10	00.	1/2% Wat	er	
slowly while stirri	ng vig	or-	Triethanolamine	-				00
ously.*			Water		ee.	Triethanolamine Water	175	00.
No. 2			Stearic Acid		cc.	Stearic Acid		
Triethanolamine		cc.	Alcohol		cc.	Alcohol	1400	ee.
Water	1	cc.	Gasoline		cc.	Gasoline	31500	
Oleic Acid		cc.	No. 11	40			0.1000	C.C.
Butanoi		cc.		_		No. 19		
Gasoline	45	cc.	Triethanolamine	1	cc.	1/4% Wat	er	
No. 3			Water	1	cc.		175	ce.
Triethanolamine	16	ee.	Stearic Acid Alcohol	2	ec.	Water	85	
Water	1				ec.	Stearic Acid	1400	ec.
Oleic Acid	1/2		Gasoline		cc.		1400	ec.
Butanol	5	cc.	No. 12			Gasoline	31500	
Gasoline	1½ 5 45	ce.	Triethanolamine					
No. 4			Water	1	ce.	No. 20		
	4.1		Stearic Acid Alcohol	3	cc.	10% Wat	ter	
Triethanolamine	1/2		Alcohol	. 2	cc.	Trihydroxyethyl-		
Oleic Acid	1	cc.	Gasoline	45	cc.	amine Laurate	3500	cc.
Water Butanol Gasoline	ī	ee.	No. 13			Gasoline Butanol Water	31500	cc.
Butanol	.e	ec.	Triethanolamine	1/2	cc.	Butanol	5600	ee.
	40	ec.	Water	1	cc.	Water	3500	
No. 5			Stearic Acid	3	cc.	Triethanolamine	1750	ec.
Triethanolamine		ee.	Alcohol	3	cc.	No. 21		
Water Oleic Acid		ec.	Gasoline	45	cc.			
Oleic Acid		cc.	No. 14			5% Wate	er	
Dutanoi		ec.	Triethanolamine	1/4	cc.	Trihydroxyethyl	0.00	
Alcohol	4	cc.	Water	1/2	cc.	amine Laurate	2100	ec.
No. 6			Stearic Acid	3	cc.	Gasoline	31500	cc.
Triethanolamine	1	ec.	Alcohol	3	cc.	Butanol Water	3500	
Water		cc.	Gasoline	45	cc.		1750	
Oloic Acid	1	cc.	No. 15			Triethanolamine	1090	ee.
Butanol Alcohol	4	cc.	Triethanolamine	1/,	cc.	No. 22		
Alcohol	2	ec.	Water		cc.	1% Wate	er	
No. 7			Stearic Acid	2	cc.	Trihydroxyethyl-		
Triothonolomina	1	ec.	Alcohol	2	cc.	amine Linoleat	0 2100	00
Water	. 1	00.	Gasoline	45	cc.	Gasoline	31500	
Oleic Acid	1	00.	No. 16			Ruterol	3500	00
Water Oleic Acid Butanol Kerosene Gasoline No. 8	5	ee.	100 10			Butanol Water	700	cc.
Kerosene	20	CC.	1% Wat			Triethanolamine	1050	cc.
Gasoline	25	cc.	Triethanolamine	175	cc.	# 1 TO CLEWING WILLIAM	2000	300
NT- 0		J.						
			Stearic Acid Alcohol	1400	cc.			
Triethanolamine		cc.	Alconol	21500	cc.			
Water		cc.						
* The stability of a colloid mill.	t the a	pove	emulsions is improve	ed cons	idera	DIY II they are pass	sed thro	ugh

Bright Drying Wax Emulsion

Paraffin Wax	15	g.
Oleic Acid	15	g.
Triethanolamine	20	ğ.
Borax) previously	$7\frac{1}{2}$	g.
Water (dissolved	$7\frac{1}{2}$	g.

Warm together to 90° C, and mix with an electric mixer. While keeping at 90–100° C, and stirring vigorously add the following which must be at 90–95° C.

9		
Carnauba Wax	100	g.
Water	1000	

Cool quickly and package.

Paraffin Wax Emulsion Formula No. 1

Paraffin	Wax	120	g.
Stearic	Acid	12	g.

Melt together and while stirring vigorously add following heated to 55° C.

Ammonia (26° Bé.) 6 cc. Water 182 cc.

Stir until uniform.

Water (boiling)

No. 2

Glyceryl	Monostearate	5	g.
Water		150	

Heat and stir vigorously until uniform. Pour into this slowly while stirring strongly:

Paraffin Wax (melted) 40 g.

Paraffin Wax Emulsion (Non-Alkaline)

(
Paraffin Wax	25 g.
Glycol Stearate	5 g.

Melt together and while stirring vigorously add

175 ec.

Laundry Calendering Wax Emulsion Mix 33 parts of paraffin wax with 3 parts of oleine, and pour this mixture into a solution of 0.6 part of strong ammonia in 63.4 parts of water, heated to 160° F.

Aqueous Fat-Dissolving Emulsion German Patent 598,216

Prepare: Carragheen Moss Dispersion, warming gently in water, remove, thicken components in a centrifugal.

Acidify with oxalic acid. Mix thoroughly.

Acidified Carragheen Moss

Solution 100 cc.
Phosphoric Acid (Free from Arsenic) (67%) 10 cc.

Fat Solvent	200 cc.
e.g. Trichloroethylene Naphtha	100 cc. 200 cc.

Chlorinated Naphthalene Emulsion British Patent 413,756

Eighty g. of wax-like chlorinated naphthalene, of setting point 93° C. is dissolved in 100 g. trichloroethylene, and is added with stirring to a warm mixture of water 60 g., Turkey-red oil 10 g., casein 3 g. and strong ammonium hydroxide 1 g.

Emulsions of Oils, Fats, Waxes and Resins

British Patent 431,642

Water is dispersed in oils, fats, waxes, resins, artificial resins, pitches, asphalts or the like by adding to the water, prior to or during the mixing, about 0.01% of the principal substance of aqueous alkali, such as caustic soda or potash or ammonia, having dissolved therein aromatic hydrocarbon derivatives or their salts soluble in alkali such as benzoic acid, sodium salicylate, o, m or p-cresol, xylenol, guaicol, or cresol, or mixtures thereof. The products may have pigments or solid substances incorporated therewith for use as paints, color varnishes, printing inks, or lubricants.

Formula No. 1

600 g. of water, containing 0.012 g. of a solution of 15% caustic soda and 1.5% of the above specified substances, are stirred at 25–20° C. into 1000 g. of linseed-oil varnish; 280 g. of the resulting water-in-oil emulsion are stirred with 530 g. of red lead and 175 g. of calcite.

No. 2

250 g. of water containing 0.02 g. of emulsifier as in (1) are stirred into a mixture of 100 g. of asphalt and 900 g. of printers' linseed-oil varnish; 9 g. of nigrosin are stirred into the product to produce a printers' ink.

No. 3

350 g. of water containing 0.015 g. of caustic soda and 0.0015 g. of sodium benzoate are stirred at 30° C. into 1000 g. of olive oil; the product may be used as salad oil.

No. 4

300 g. of water containing 0.02 g. of caustic potash and 0.02 g. of cresol are stirred into 1000 g. of viscous mineral lubricating oil and 100-200 g. of

graphite added with stirring to produce a lubricant.

Emulsions of Oils, Fats, or Waxes German Patent 575,922

Formula No. 1

Cod Liver Oil Pectin Milk Sugar		80 0.5 20	g g g
Water	No. 2	20	g.
Paraffin Oil Pectin Milk Sugar Water		80 0.5 20 20	80 80 80 80

German Patent 585,586, Addition to the Above (575,922)

For stable emulsions containing up to 80% of oils use instead of Milk-Sugar:

Fruit Sugar (Fructose) Invert Sugar (Invertose) Grape Sugar (Glucose) Manna Sugar (Mannose) Never use Cane Sugar!

Pine Oil Emulsion

Pine Oil	9 g.
Diglycol Laurate	4 g.
Mineral Oil	1 g.
Water	100 g.

Mix first three materials, and then add water slowly while stirring vigorously.

Cottonseed Oil Emulsion

Diglycol Laurate	18 cc.
Cottonseed Oil	40 cc.
Water	50 cc.

China Wood Oil Emulsion

Diglycol Laurate	18	cc.
China Wood Oil	40	cc.
Water	55	cc.

Mineral Oil Emulsion Cream

Glyceryl	Monostearate	5 g.
Water		125 ec.

Heat together and stir until uniform then add slowly while stirring vigorously Mineral Oil 43 cc.

Soluble Oil

U. S. Patent 1,965,935

A soluble oil composed of the following ingredients has unique emulsification and stability properties:

Sodium	Corn	Oil	Soap	14	g.
Water				6	g.

Mineral Oil Water White Rosin	$\frac{64}{10}$	
"Carbitol" (Monoethyl	0	

Ether of Diethylene Glycol) 2 g. Diethylene Glycol 4 g.

This oil is clear and will not become cloudy when cooled to a temperature of 60° F. and will not become covered with a film after standing exposed to the air at a temperature of 80° F. over a long period of time or at a temperature of 200° F. for one day. This oil will readily emulsify with water after standing exposed to the air at 200° F. for two days. Aqueous emulsions containing this oil are very stable even at a temperature of 200° F. In general, stable aqueous emulsions are prepared by using 1% to 35% of this oil, although stable aqueous emulsions can be prepared by using proportions of the oil outside these limits.

Carbon Tetrachloride and Tetrachloroethylene Emulsions

The following formula may be used for a 50% preparation: 20 g. of commercial soft soap, 6 cc. of cresol, 50 cc. of carbon tetrachloride or tetrachloroethylene and 100 cc. of liquid paraffin.

Phenol-Formaldehyde Resin Emulsion Australian Patent 17,583

Phenol-Formaldehyde Resin	45	g.
Paraffin Oil	5	g.
(Heat together) Sulphonated Sperm Oil		-
Sulphonated Sperm Oil	5	g.
Olein	5	g.
Cyclohexanol	1	g.

Partially saponify above with aqueous caustic soda then add

Glue			2	1/2 g.
Water			45	g.
Mix in	homogenizer	or	colloid	mill.

Synthetic Resin Emulsion U. S. Patent 1,999,715

One hundred parts, by weight, of ground resorcinol are placed in a metal container or kettle which is jacketed with an outside container, in turn equipped for steam heating and water cooling. This permits of temperature regulation during the chemical reaction as it is very essential to carefully control the temperature of the reacting substances in order to prevent their conversion to the insoluble

stage. The kettle may be equipped with suitable agitators to permit the rapid churning of the kettle contents. To the resorcinol, 112 parts, by weight, of 37.5% formaldehyde solution (formalin) are added, and the temperature is increased to a maximum of 60° C., so that

the resorcinol will dissolve.

In a separate container, 8 parts, by weight, of para-nitraniline are dissolved in 20 parts, by weight, of cresol having a boiling point range of from about 215° C. to 230° C. The melted paranitraniline is added to the formaldehyde solution, and the mass is thoroughly agitated, the reaction temperature being raised to from 70° C. to about 75° C. A plasticizer of vegetable or animal oils and wax with a filler and suitable coloring material may be added as a paste to the mixture in the kettle. As an example, employ 8 parts, by weight, of clay, 0.8 part, by weight, of beeswax, and 1 part, by weight, of iron oxide, all best ground in a paint mill or ball mill to obtain thorough dispersion. The paste has been found to mix readily with the thickening

liquid in the kettle.

For best results, the temperature should be maintained at no higher than 80° C. When the mass in the kettle becomes stringy, and before gelatinization can take place, the emulsification enhancing agent is added. A water solution (65 parts, by weight) containing 0.1% of borax is added, first slowly, and then rapidly to the agitated resin. The borax, or any other suitable alkaline salt, serves to so enhance the emulsifying action of the wax and/or the oils (and gums, if any are used) as to enable them to properly maintain the resin in suspension in the water. This results from the fact that the borax or other alkaline salt used reduces the surface tension of the water from its normal value at the operating temperatures, thereby more readily effecting a wetting of the resin particles and enabling the wax or other fatty material used to more easily retain the resin particles in suspension. The temperature is dropped to about 20° C. and thickening of the resin takes place. A solution of alcohol (about 65 parts, by weight) or an equal quantity of a 20% benzol or toluene solution in alcohol may be added to the emulsifier to obtain proper consistency. The varnish is immediately strained to remove any foreign particles. The mass is then cooled, with accompanying increase in viscosity of the varnish, and additional alcohol or solvent mixture may be added to obtain the desired viscosity.

Chlorinated Rubber (Tornesit) Emulsions

U. S. Patent 2,008,558

Formula No. 1

50 parts, by weight, of toluene, 50 parts of water and 20 parts of pulverulent chlorinated rubber are introduced jointly into a stirring apparatus and stirred. A uniform and stable dispersion is formed in a few minutes.

50 parts, by weight, of toluene and 50 of water are brought together and intimately stirred, 20 parts of finely divided solid chlorinated rubber being added during the stirring operation. A uniform and stable dispersion is formed immediately.

Chlorinated Rubber Emulsion British Patent 414,072

20 parts, by weight, of oleic acid, saponified with 20 parts of sodium silicate in 200 parts of water, is stirred at 100° C. into 5 parts of chlorinated rubber dissolved in 25 parts resin oil; 125 parts of water containing casein 8 and ammonia 0.5 parts is then added.

Aqueous Dispersions of Bitumen German Patent 557,228

Soya Bean Meal		1	g.
a. Soya Bean Meal Water	4	9	cc.
b. Caustic Soda		0.2	cc.
c. Bitumen Mixture,			
Liquefied		ś٨	œ

Boil a after soaking, then saponify with b and emulsify c, stirring vigorously.

Tar Emulsion Austrian Patent 137,894

Crude	Montan Wax	3	lb.
Crude	Wool Fat	2	lb.
Tar		95	lb.

Heat to 80-90° C. and while mixing vigorously run into a 1% caustic soda solution heated to 60° C.

Asphalt Emulsion U. S. Patent 1,931,072

An aqueous solution of soap (9 parts) by weight, is dissolved in warm water (78 parts), and a low grade fuel oil or crude asphaltic-base oil (20 parts) is dispersed therein. A relatively small quantity (1-2 parts) of a metallic salt of a fatty acid, e.g., aluminum oleate is mixed therewith, the emulsion is warmed, and asphalt (296 parts) is added slowly and with agitation, and distributed uniformly throughout the mixture.

Non-Rusting Alkaline Emulsions Latherless Shaving Cream U. S. Patent 1,968,722

Stearic Acid	22 lb.
Glycerin	10 lb.
Ammonia (28%)	1 lb.
Sodium Nitrite	0.1 lb.
Water	67 lb.

The stearic acid is first heated to about The glycerin and water are then mixed together apart from the stearic acid and also heated to about 85° C. To the glycerin and water add the ammonia. This solution is then poured into the stearic acid and thoroughly stirred. When the whole mix is cooled add sodium nitrite.

Polish

Carnauba Wax	12	g.
Rosin	0.5	g.
Triethanolamine Oleate	3.5	g.
Sodium Nitrite	0.1	g.
Water sufficient to make	100	g.

Library Paste

Starch	24 g.
Gum Acacia	3 g.
Glycerin	6 g.
Borax	0.5 g.
Sodium Nitrite	0.1 g.
Oil of Cloves	0.1 g.
Water	72 g.

Soap Base Lubricating Emulsion Cottonseed Oil 3 kg. Mineral Oil 1-2 kg. 132 g. Caustic Soda

Heat to 180° C. until foaming stops. Add 13 kg. mineral oil in successive portions at intervals with stirring bringing up to 190-210° C. Pour into wooden tubs and cool to 70° C. Add 9 kg. water with stirring.

High-Molecular Organic Sulphonic Acid Emulsifier

German Patent 616,321

Formula No. 1

Yellow Oil from Brown	
Coal Tar	100 g.
Paraldehyde	10 g.
Chlorosulphonic Acid	125 g.
Add the acid at 30-35° C	cool, stir

sulphonic acid from unchanged oil. Pour into 3 parts of ice-water neutralized with concentrate caustic soda. Let stand, separate from impurities, dry in vacuum.

No. 2

Same, but substitute Paraldehyde by Acetalde- hyde (50%)	20 g.
No. 3	
As No. 1, but use	
Paraffin Óil from Brown	
Coal Tar	100 g.
No. 4	
Solar Oil from Brown	
Coal Tar	100 g.
Heptaldehyde (Oenanthol)	20 g.
Chlorosulphonic Acid	150 g.
At 35° C. has to stand 1	day, other
act as in No. 1.	
No. 5	

Paraffin Oil (7.5° E at	
20°)	100 g.
Benzaldehyde	15 g.
Chlorosulphonic Acid	130 g.
Method as No. 1	

Sulphonation of Cetyl Alcohol

Melt the		
Cetyl Alcohol	40	g.
Dissolve in		_
Acetic Anhydride	20	g.
Treat with		_
Sulphuric Acid (Concentrated		
or Fuming)	40	g.
The reaction is run below 10°	C.	

Sulphonating Napthenic Alcohols U. S. Patent 2,000,994

One part by weight of a raw commercial naphthenic acid (boiling point 90-230° C. at 13 mm. pressure) is dissolved in 2 parts by weight of 3% butyl alco-holic hydrochloric acid and heated to boiling for four hours. The butanol and hydrochloric acid are then distilled off and 200 kg. of the naphthenic acid so treated are reduced in an autoclave with 90 kg. of sodium and 1,000 kg. of butyl alcohol. The whole is then heated under constant agitation to 140° C. for 1½ hours. After cooling to 90° C. the reaction mass is poured into water, the underlying liquor is drawn off and the remainder is neutralized and washed several times. It is then dried over lime and the excess butyl alcohol is removed by distillation. The product so obtained boils between 70 and 230° C. at 10 mm. thoroughly. After 18 hours separate the | pressure and posseses an acetyl saponification number 175 and an iodine number 22. It is free from saponifiable components and dissolves to give a clear solution in concentrated sulphuric acid. Dilution with water produces no turbidity. The conversion of the product into the sulphuric acid derivative can be carried

out in the following manner:

20 parts by weight of chlorosulphonic acid are gradually added to 50 parts of the above mentioned product and to this are subsequently added 5 parts of sulphuric acid whereupon the temperature rises to 40° C. The reaction mass is then washed with salt solution and neutralized. Upon evaporation in vacuo the sulphonate and/or sulphate is obtained in a solid grindable form.

Sulphonating Oils

A. Cod, Sperm, Cottonseed and Castor
Oils

1. High Sulphonation Product

Any of the Above Oils 735 lb. Sulphuric Acid 275 lb.

Run in the acid in a thin jet as quickly as possible with good mixing but do not allow temperature to rise above 95° F. Agitate for 5 or 6 hours until a sample in the case of cod oil is soluble in distilled water without opalescence. With cottonseed oil the solution will be slightly translucent. The oil is now dropped into the mixing tank, containing two and onehalf times the volume of oil of Glauber's salt solution, 10° Bé. Agitate smoothly for five to ten minutes and warm to 104° F. Allow to separate. Draw off the water and make the oil nearly neutral to methyl orange with caustic soda. Allow to stand over night. It is to be noted. that according to the acidity of the oil at this stage, when allowed to stand, the amount of free fatty acid in the finished oil can be regulated. Next morning, draw off the water again, and clear with caustic soda.

B. Red, Cod, Castor, Neatsfoot or Refined Corn Oils

2. Quick Sulphonation Method

Any of the Above Oils 775 lb. Sulphuric Acid 225 lb.

Usually used for cleic acid, cod oil, castor oil, Neatsfoot oil, refined corn oil, and mixed oils. Sulphuric acid $=22\frac{1}{2}\%$ on the weight of the oil.

The acid is run into the oil quickly while the oil is violently agitated. With a ten-barrel batch of oil, the acid takes about thirty minutes to run in. The temperature rises quickly and as soon as it

reaches 130-135° F., the oil mixture is dumped quickly into a mixing tank situated underneath the sulphonating tank. The mixing tank contains Glauber's salt solution 10° Bé. equal to double the volume of the oil. The Glauber's salt solution is at room temperature. The oil and Glauber's salt solution is agitated smoothly for five to ten minutes and the oil allowed to separate. Separation is nearly complete in half to one hour. The clear water is drawn off to a storage tank, and after neutralizing with caustic soda, is used over again for the next The oil is neutralized with caustic soda until it is nearly neutral to methyl orange, that is, slightly on the acid side. Allow the oil to stand until morning and a further separation will take place. When the oil is completely separated, and the water drawn off, the oil should test 20% water. It is now cleared by the addition of further caustic soda. In winter time, it is better to use caustic potash for the final finishing, as it gives a more liquid oil. In testing the acidity of the oil, after the first separation, it is recommended to use an ether and salt solution for the titration with methyl orange.

C. Castor Oil, Concentrated

Castor Oil 1000 lb. Sulphuric Acid (100%) 1000 lb.

Dilute the castor oil with ethylene dichloride. Run the acid in slowly to the previously cooled oil and solvent mixture. Do not allow the temperature to rise above 60° F. After the acid is all in, continue stirring until a few drops dissolve perfectly clear in distilled water, and also dissolve perfectly clear in a saturated solution of calcium sulphate. Do not continue stirring after this point, but then add to it a 5% solution of Glauber's salt solution, equal in volume to three times that of the sulphonated mixture. The solution of Glauber's salt is kept cool by means of ice. The temperature not being allowed to rise above 60° F. Allow to separate, and wash twice with 25% Glauber's salt solution. Separate, and add caustic soda until neutral, and then distil off the solvent.

D. Oleic Acid and Ricinoleic Acids, Concentrated

Above Fatty Acids 100 lb. Sulphuric Acid (100%) 100 lb.

On a large scale some manufacturers use a dough type mixer, brine cooled, while others use a system, wherein the sulphuric acid and oil are sprayed simultaneously by a whirl disc system into a

large reaction vessel, being sufficiently cooled previously so that the heat of reaction does not cause the product formed to become unduly heated before running out of reaction vessel. Sulphonation uses 100 lb. fatty acids and 100 lb. sulphuric

acid 100 per cent strength.

Keep temperature below 50° F. while adding the sulphuric acid. Sulphonation time is 50-60 minutes. Wash with Glauber's salt solution 12-15° Bć. twice, keeping temperature below 70° F. Let stand over night to separate and neutralize with caustic soda. The product is allowed to stand 3-5 days at 15-20° C. to allow the Glauber's salt to crystallize out. This crystallization can be improved by the addition to the oil of a small quantity of a volatile solvent such as xylene, trichloroethylene, carbon tetrachloride, etc.

Sulphonation of Castor Oil French Patent 745,787

a. Castor Oilb. Sulphuric Acid (66° Bé.)100 kg.

Add b to a in very thin jet (2 hours) and with continuous stirring, keeping the temperature at 10-13° C. Wash product once or twice with salt-solution, keeping the temperature below 15° C. Separate oil from the aqueous layer in the usual way, neutralize.

Sulphonating Oils U. S. Patent 1,967,655

Formula No. 1

100 kg. of ricinoleic acid are sulphonated at temperatures below 5° C. with 90 kg. of 30% oleum. 30 kg. of glycol mono-methyl ether are added to the crude sulphonation product. After completion of the reaction, ice is added and the product washed with Glauber's salt solution.

No. 2

100 kg. of 12-hydroxy stearic acid are mixed with 65 kg. of glycol mono-ethyl ether and sulphonation effected at temperatures below 0° C. with 36 kg. of chlorosulphonic acid. The product is worked up as in No. 1.

No. 3

100 kg. of naphthoic acid are sulphonated with 70 kg. of chlorosulphonic acid, 55 kg. of glycol mono-methyl ether are added to the crude sulphonation product. In place of sulphuric acid and the like sulphonating agents, alkyl sulphuric acids

or alkyl chlorosulphonic acids may be employed.

Sulphonation of Fatty Oils, Fats, Waxes Austrian Patent 134,993

 $\begin{array}{ccc} \text{Whale Oil (Sperm Oil)} & 1 \text{ lb.} \\ \text{Spindle Oil} & 3-4 \text{ lb.} \\ \text{Fuming Sulphuric Acid} \\ (30\% \text{ SO}_3) & 1 \text{ lb.} \end{array}$

Run reaction at 40-45° C., adding sulphuric acid in a jet. Stir, then let stand 12 to 24 hours; wash with sodium chloride- or sodium sulphate-solution, separate from acid wash water. Neutralize, if necessary, with organic bases, until a drop of oil, when diluted with water, shows nearly no turbidity.

Cellulose Ester Emulsions U. S. Patent 1,970,572

A pyroxylin base is prepared by colloiding 12.5 parts by weight of alcoholwetted pyroxylin (10 parts of dry ½" pyroxylin) with 20 parts by weight of blown linseed oil in a suitable mixer, such as the Werner and Pfleiderer mixer. 25 parts by weight of a solvent mixture are then added to the colloided mass in portions equalling 5 parts by weight to form a homogeneous base having the following composition:

Pyroxylin (½ sec.) 10 g.
Alcohol (Denatured) 2.5 g.
Blown Linseed Oil 20 g.
Butyl Acetate 20 g.
Butyl Lactate 5 g.

An emulsion is prepared by heating 0.5 part by weight of sodium oleate with 15 parts by weight of gasoline to a clear gel; after which 2 parts by weight of water are added to the hot gel with vigorous stirring, thus forming a concentrated emulsion of gasoline in water that is stabilized by sodium oleate. For convenience this will hereafter be called the agent emulsion.

The presolution or solvating of the sodium oleate in gasoline or some similar liquid is desirable to assure uniform dis-

tribution.

17.5 parts by weight of the agent enulsion are then stirred vigorously into 57.5 parts by weight of the pyroxylin base with a high speed stirrer of the propeller blade type.

propeller blade type.

Inversion of the emulsion from the

water-in-oil type to the oil-in-water type may be effected in various ways, as explained below, but in this example it is effected by the sudden addition of water in relatively large quantities, the time of

addition being the controlling factor in particle size, as indicated by systems a, \bar{b} , and c.

System (a): 20 parts by weight of water are added in small portions with vigorous stirring, thus yielding a viscous water-in-oil type dispersion. 10 parts by weight of water are then added with vigorous stirring to invert the system to the oil-in-water type. 68 parts by weight of water are added next, either slowly or rapidly, with more moderate stirring. Microscopic measurements of particle size average 1.19 microns, and the dispersion spontaneously wets an absorbent

type of paper.

System (b): 35 parts by weight of water are added in small portions with vigorous stirring, yielding a viscous water-in-oil type dispersion. 10 parts by weight of water are then added with vigorous stirring to invert the system to the oil-in-water type. 53 parts by weight of water are next added, either slowly or rapidly, with more moderate stirring. The average particle size is 1.92 microns, and the dispersion does not wet paper spontaneously. Vacuum filtration is required in order to effect paper penetration, and some separation of disperse phase occurs on the surface of the paper.

System (c): 90 parts by weight of water are added in small portions with vigorous stirring, yielding a viscous water-in-oil type dispersion. 8 parts by weight of water are then added with vigorous stirring to invert the system to the oil-in-water type. The average particle size is 2.23 microns. Severe separation of the disperse phase occurs on the surface of the paper during vacuum filtration, and the dispersion is not adapted to paper impregnation.

Petroleum Demulsifier

Diglycol Laurate	83	g.
Sodium Silicate	5	g.
Rosin Soap	5	g.
Phenol	4	g.
Water	11/2	g.
Paraffin	11/2	g.

Margarine Emulsifier

Refined and deodorized sunflower oil oxidized with a current of dry air at 250° C. for 10 hours shows better emulsifying properties than Paalsgaard oil. When 0.4% of this oil is added to the emulsified mixture in the manufacture of margarine, the product after standing 54 hours retains the good taste and odor and high moisture content (14.6%).

Breaking Petroleum Emulsions U. S. Patent 1,976,602

React 250 lb. of phthalic anhydride with 500 lb. of castor oil at a temperature of approximately 120 to 145° C. for approximately 6 to 12 hours. The reaction can be followed roughly by withdrawing a small sample of the partially reacted mass and permitting it to cool on a watch glass. When the reaction is completed, crystals of phthalic anhydride no longer When the sample no longer shows the presence of such crystals on cooling, it can be titrated with a standard volumetric alkaline solution, so as to indicate that the acid which remains is due entirely to the carboxylic hydrogen and not due to any unreacted phthalic anhydride. One must guard against a rise in temperature.

The product of reaction represents a viscous yellow oil not unlike blown castor oil in consistency. It is neutralized with sufficient ammonium hydroxide to completely convert all acidic material into the ammonium salt. The product thus obtained is substantially water-soluble

and is suitable for use.

A treating agent or demulsifying agent of the kind described may be brought in contact with the emulsion to he treated in any of the numerous ways now employed in the treatment of petroleum emulsions of the water-in-oil type with chemical demulsifying agents, such for example, as by introducing the treating agent into the well in which the emulsion is produced, introducing the treating agent into a conduit through which the emulsion is stored, or introducing the treating agent into a container that holds a sludge obtained from the bottom of an oil storage tank. In some instances, it may be advisable to introduce the treating agent into a producing well in such a way that it will become mixed with water and oil that are emerging from the surrounding strata, before said water and oil enter the barrel of the well pump or the tubing up through which said water and oil flow to the surface of the ground. After treatment the emulsion is allowed to stand in a quiescent state, usually in a settling tank, at a temperature varying from atmospheric temperature to about 200° F., so as to permit the water or brine to separate from the oil, it being preferable to keep the temperature low enough so as to prevent the valuable constituents of the oil from volatilizing. If desired, the treated emulsion may be acted upon by one or the other of various kinds of apparatus now used in the operation of breaking petroleum emulsions, such as homogenizers, hay tanks, gun barrels, filters, centrifuges or electrical dehydrators.

The amount of treating agent on the anhydrous basis that is required to break the emulsion may vary from approximately 1 part of treating agent to 500 parts of emulsion, up to a ratio of 1 part of treating agent to 20,000 parts of emulsion, depending upon the type or kind of emulsion being treated. In treating exceptionally refractory emulsions of the kind commonly referred to as "tank

bottoms" or "residual pit oils," the minimum ratio above referred to is often necessary, but in treating fresh emulsions, i.e., emulsions that will yield readily to the action of chemical demulsifying agents, the maximum ratio above mentioned will frequently produce highly satisfactory results. For the average petroleum emulsion of the water-in-oil type a ratio of 1 part of treating agent to 10,000 parts of emulsion will usually be found to produce commercially satisfactory results.

FARM AND GARDEN SPECIALTIES

Tree	Bands	for	Caterpi	llar	and	Flies	
		Horn	oM elm	1			

a. Rosin Oil Spindle Oil	9 g.
a. Spindle Oil	20 g.
Slaked Lime	6-9 g.
b. Slaked Lime Spindle Oil	65-62 g.

Add a to b, stir violently to homogenize. Stir until congealing starts. Allow to set for 24 hours.

No. 2

Rosin	30	g.
Linseed Oil* Varnish	20	ğ.
Beeswax, Yellow	2	g.

* Or Rape Seed Oil, or Wool Fat, when a longer catching period is desired.

The melted and well mixed glue is put on the bark of the tree; over it put a ring of cloth, fastened with wire, then put over that again a layer of glue, all around the stock.

No. 3

Colophony Linseed Oil Varnish	300 200	
Yellow. Wax	20	g.

Protecting Mixture for Young Trees Against Game

Ceresin (58-60° C.)	20	oz.
Spindle Oil, Distilled	60	oz.
Dippel's Animal Oil or		
Carholineum	20	OZ

Melt up and stir until cold.

Codling Moth Tree Bands Formula No. 1

Cloth is impregnated with

Beta Naphthol Crude 1 lb.
Red Engine Oil 1.5 pt.

Apply at 130–132° F.

No. 2

Beta Naphthol Crude	1	lb.
Mineral Oil (200-300 sec.)	11/2	pt.
Gasoline	1	pt.

210.0		
Water	2	gal.
Ammonia (28%)	2.4	fl. oz.
Casein	4	OZ.
Mineral Oil, Refined		

(65-75 sec.)

Grafting Wax Formula No. 1

	Colophony Beeswax	350 g.
	Beeswax	10 g.
a.	Pitch	60 g.
	Linseed Oil	25 g.
	Linseed Oil Turpentine, Venice	15 g.
b.	Methanol	85 g.
M	elt up a, then stir until	cool, add b.

No. 2

Linseed Oil Turpentine Oil	1	lb.
Turpentine Oil	4	lb.
". Beeswax	3	lb.
Colophony	9	lb.
b. Methanol	1	lb.
Dissolve a continualy thin	with h	

No. 3

110. 0			
Castor Oil	1/4 lb.		
Rosin	5 lb.		
Beeswax	1 lb.		
Charcoal	34 lb.		
Glyceryl Monostearate	1 lb.		

Melt and apply with brush. This excludes air and fungi; prevents drying out and doesn't injure live tissues.

Bleaching Citrus Fruit Blemish

Navel oranges, badly blemished with sooty blotch, are thoroughly cleaned by dipping for ½-1 min. in a solution containing 0.25 lb. each of boric acid and chloride of lime.

Removing Arsenic Residues from Fruits Wash with a 1% solution of ammonia or caustic soda.

Preserving Color of Leaves

Immerse leaves in	
Glycerin	5 g.
Copper Sulphate	2 g.
Water	93 cc.

Non-Poisonous Fly-Papers

TOM TOROUGH	rij rapeis		
Quassia	16	oz.	
Colocynth	2	oz.	
Long Pepper	4	OZ.	
Water	to make 1	gal.	
20 12 142 12 2		_	

Boil until the decoction is reduced to 4 pints; strain; dissolve in the clear

gal.

liquid 4 oz. of sugar. Dip the absorbent paper in this solution.

Cobalt Fly-Papers

Dissolve cobalt chloride, 1 oz., and Tartar Emetic, 1 d., in 1 gal. of the Quassia decoction (formula above), and dip the paper in the resulting solution.

Fly Catcher		
Colophony (Rosin) G	49	g.
Mineral Oil (Viscosity		
3½-4° E at 50° C.)	36	
Lanolin, Anhydrous	4	g.
Beeswax, Pure	1	g.
Castor Oil	2	g.

Moth Powder

Camphor	4	oz.
Benzoin	1	0 Z .
Black Pepper	2	$0Z_{\bullet}$
Cedar Sawdust	5	oz.

Mix after reducing the solids to a coarse powder.

Roach Eradicating	Powder	
Sodium Fluoride	60	oz.
Wheat Flour	20	oz.
Corn Starch	12	oz.
Cocoa	8	oz.

The sodium fluoride should be in a finely powdered form and thoroughly mixed and then sifted to make certain of a homogeneous product. This may be made into a paste with a minimum of water and placed in new or used crown caps, allowed to dry and laid in roach infested places. It may also be dusted as a powder. The filled caps, however, can be used over again and are cleaned up more readily than the powder.

Mosquito Spray for Outdoor	Gathering
Kerosene Containing Py-	
rethrum Extract Equiva-	
lent to 1 lb. of Flowers (Analyzing 0.9% Pyreth	
rins) per Gallon	100 gal. 50 gal.
Water	50 gal.
Sodium Laurel Sulphate	
(Emulsifier)	6 lb.

The emulsifier is first mixed with the water and transferred to the tank. The oil is then run in gradually into the tank with agitators and pump working at full speed. After all the oil has been added the pumping is continued until the entire mixture has passed through the hose and

back into the tank two or three times or until the mixture is thick and homogeneous, showing no free oil on the surface. The finished product is then pumped into drums for storing. This constitutes the stock emulsion. Excessive foaming may be eliminated by dissolving about two or three pounds of wool grease (degras) in the kerosene before emulsifying. Any other suitable apparatus for emulsification can be used.

The cost of preparing the concentrated emulsion is about 23 cents per gallon, based on the present price of pyrethrum, which makes out slightly over 2 cents per spray gallon. When purchased, the stock emulsion costs from 30 to 50 cents per gallon, depending on the quantity ordered.

Directions for Spraying

About half an hour before the gathering takes place the area is completely sprayed with the larvacide diluted 1:10 or 1:12, that is 1 part of larvacide is mixed with 10 or 12 parts of water. The spraying is done with a power sprayer capable of developing a pressure of 100 pounds or more per square inch and equipped with a spray gun. Before mixing with water the concentrated stock larvacide should be well shaken. Also the diluted spray should be frequently stirred or agitated in order to secure uniform distribution throughout the spraying operation. The spray is applied in the form of a fine fog directly to the grass, grounds, tents, trees, shrubs, etc. Then the stream is directed upward so as to saturate the atmosphere with the fog. At no time should a coarse spray be applied, since it is unnecessary and may injure vegetation. The grounds for about 20 feet outside the area should also be thoroughly fogged, especially when tall grass, shrubs, woodland and other vegetation are present offering a hiding place from which adult female mosquitoes may issue suddenly at dusk in large numbers. If the area has been thoroughly fogged one treatment may suffice for two hours or even the rest of the evening. If mosquitoes become bothersome later in the evening, the area on the outside of the "gathering" grounds should again be fogged, directing the stream primarily upward and towards the ground to be protected. This outside fogging may be repeated again if necessary. On small areas, such as back-yards, private lawns, etc., a knapsack sprayer or bucket pump capable of producing a fog spray, of 10 to 15 feet high, can be used.

Weed Killers Formula No. 1

Poisonous:

Arsenite of Soda (Concen-

trated Solution) 1 gal. 20 gal. Water

Mix through and sprinkle on vegetation to be exterminated, making application on a bright clear day.

Non-Poisonous:

1 lb. Chlorate of Soda Water 1 gal.

Dissolve chlorate of soda in the water and use this solution without further dilution by sprinkling on vegetation wished exterminated.

Weed Killer British Patent 418,061

Formula No. 1

Ammonium Chloride Copper Sulphate Calcium Carbonate	83 g. 5 g. 12 g.
No. 2	
Asymanium Chlorida	95 ~

Ammonium Chloride 25 g. 25 g. Sodium Nitrate 50 g. Ferrous Sulphate

Ragwort Weed Killer

Use ammonium sulphocyanide (5-10% solution), 200 gal. per acre. weather is best time.

Killing Weeds on Lawns

To kill weeds on lawns, golf courses, etc., treatment with a solution of ammonium sulphate and soft soap has been found effective. A mixture adopted for this purpose in England contains 1 lb. of ammonium sulphate, 1/2 lb. of soft soap, and 1 gal. of water, to be used for every 8 square yards.

Hydrogen Sulphide as an Insecticide and Fungicide

Extensive trials carried out by the Azov-Black Sea branch of the All-Union Institute of Plant Protection have proved that hydrogen sulphide may be successfully used for rodents, insects, and fungi Laboratory experiments have shown that a 0.02 to 0.03% concentration of hydrogen sulphide in air is sufficient to kill the earless marmot. Field experiments proved that 4 to 6 g. of hydrogen sulphide per burrow are quite sufficient, while in better conditions (i.e., when the

soil is warm and dry) this rate may be reduced to 3 g. per burrow. The same results are obtained when applying sulphuric slags, which emit hydrogen sulphide because of the action of moisture absorbed from the air. In this case some 8 to 9 g. of slag per burrow are suffi-cient, the mortality of earless marmot reaching 92 to 98%.

Hydrogen sulphide proved especially efficient as a means of destroying barn mites, being more penetrable in grain than chloropicrin and carbon disulphide.

Experiments made in the huge Millerovo elevator have proved the practicability of this method. Exposure for 40 hours at a rate of 400 g. of hydrogen sulphide per ton of grain proved efficient. Hydrogen sulphide does not reduce the germination rate of seeds and only a few strains decrease their germination with 4 to 8%, while the majority of strains even increase their germination rate with 15 to 30 per cent. Experiments on feeding the treated grain to cocks and rabbits have shown that no injury has resulted.

Fair results have been obtained, too, when applying hydrogen sulphide as a fungicide. Laboratory experiments have proved that the spores of main fungous diseases of seeds perish when seeds are exposed to hydrogen sulphide for 1 to 4 days at a rate of 200 to 400 g. of gas per cu. m. (smut and bacteriosis of cereals, goummosis of cotton, bacterial rot of vegetables).

Red Squill Extract

Extract 15 g. red squill by repeated extractions with 100 cc. boiling methanol in an enclosed system percolator.

Insect Spray Formula No. 1

Petroleum Spirits	1000	cc.
Pyrethrum Extract	5	g.
Sassafras Oil	5	cc.
Methyl Salicylate	20	cc.
No. 2		
Petroleum Spirits	550	cc.
Vaseline Oil	450	cc.

Sassafras Oil 10 cc. 10 g. Pyrethrum Extract

20 cc.

Insecticide Spray Spreader

Methyl Salicylate

Water	3	5	cc.
Caustic Potash		7.4	g.
Pine Tar Oil		44.3	cc.
"Cellosolve"		10	cc.
Oleic Acid		33.3	cç.
Mix in the order given.			

Light Stable Insecticide Spray U. S. Patent 2,011,428

Gum Ghatti 2.4 lb. Cresylic Acid 0.18 lb. Water 35 lb.

White Oil (80 sec. Saybolt

at 100° F.) 62.4 lb. 1,4 Toluido Anthraquinone 0.02 lb.

The concentrated emulsion may be prepared by intimately mixing the ingredients in a colloid mill or by passing the mixture through a centrifugal pump or in any other suitable manner to give a concentrated emulsifiable composition which may readily be diluted to yield an emulsion suitable for spraying purposes.

Codling Moth Control by Nonarsenical Sprays

Sprays containing nicotine sulphate (1:640) and white oil (1:80) gives much better control of the codling moth than lead arsenate sprays.

Non-Poisoning Fruit Spray Formula No. 1

Diglycol Laurate Pyrethrum Extract	5	qt.
(20 Fold) Water No. 2	3½ 100	pt. gal.
Derris Extract (5%) Skim Milk Powder		qt. lb.

Water No. 3

Derris Root, Ground 10 lb.

90 gal.

5 lb.

Orange Worm Spray Formula No. 1

Potassium Aluminum Fluoride 50 lb. Fiber Tale 45 lb. Mineral Oil, Refined

(70 Viscosity) Use 1 lb. per tree.

Filler or Diluent

No. 2

Sodium Aluminum Fluoride 3 lb. Water 100 gal. Liquid Blood Albumin

Ŝpreader ⅓ pt.

Peach Tree Spray

A combination of the lead arsenate and zinc-lime sprays is effective not only against chewing insects such as curculio and codling moth, but against bacteriosis. The formula is:

Zinc Sulphate	8 lb.
Hydrated Lime	8 lb.

Water	100	gal.
A 44.		0

Lead Arsenate 3 lb

The spray should be used as soon as prepared.

Prune Worm Spray

Pyrethrum Extract	1	qt.
Kerosene	6	gal.
Neutral Soap	4	Ϊb.
Water	94	gal.

Pear Tree Blight Injection U. S. Patent 2,017,269

Pine Tar Oil	1	oz.
Turpentine	16	oz.

Gladiolus Thrip Spray

Manganese Arsenate	(20%)	
Arsenic)		1 lb.
Brown Sugar	60	i 1b.
Water	100	gal.

Adhesive for Hydrated Lime in Sprays

A spray of 20 lb. calcium hydroxide and 3 lb. aluminum sulphate in 100 gal. of water will give an adherent white spray residue which is repellent to the Japanese beetle. The mixture may be of value as an adherent for other spray ingredients.

Lead Arsenate Substitute

This compound is prepared by fusing 1 part diphenylamine with 2 parts sulphur at 180° C., iodine being used as catalyst. Upon recrystallizing from toluene, the light yellow crystal compound melting at 180° C., neutral, insoluble in water, slightly soluble in cold mineral oils and the usual organic solvents, is obtained. In laboratory tests, the compound is as effective as lead arsenate for codling moth larvae.

San Jose Scale Spray

Creosote	Oil	Emulsion	1	lb.
Mineral	Oil	Emulsion	3	lb.

Scale Insect Poison

Paraffin Oil	11/2	gal.
Ferrous Sulphate	(5	lh.
Caustic Soda	20	lb.
Quicklime	3	lb.
Water to make	100	gal.

Holly Sprays

Use a 3% oil emulsion containing a little nicotine sulphate. This prevents scale on living trees.

Cut holly is freed from insects by dipping and soaking for 10 minutes at 24° C. in

Derris Spray

U. S. Patent 1,934,057

Derris Extract	1-25	oz.
White Mineral Oil	40-80	oz.
Soap	5-25	oz.
Water	up to 35	oz.

The above is used diluted with water to give a mixture containing 0.06-0.25% Derris extract.

Fungi Spray U. S. Patent 2,000,843

Soft Soap	33 lb.
Cresol Soap (2% Solution)	11 lb.
Tobacco Extract (10%)	17 lb.
Potassium Permanganate	
(1/2 Normal)	22 lb.
Vegetable Glue	17 lb.
Alcohol	1/4-2 lb.

Lime, Sulphur and Salt Wash Formula No. 1 No. 2 No. 3 No. 4

Lime	2	11/2	$1\frac{3}{4}$	2	lb.
Sulphur	$1\frac{1}{2}$	11/2	$1\frac{3}{4}$	$1\frac{3}{4}$	lb.
Salt	1	11/2	134	1	lb.
Water	4	4	4	4	gal.

Boil the lime and sulphur together in a little of the water, and when combined add the rest of the water and salt. Effective as a winter application for scale.

Lime Sulphur Spray

Directions for making 50 gal. of lime sulphur spray are as follows:

apara apara, and an areas	
Sulphur	8 lb.
Spent Carbide Residue	3 gal.
Calcium Arsenate	8 oz.

Heat about 1/3 of the total amount of water, adding the sulphur slowly to make a thick paste. When the water is hot, add all the carbide residue, thoroughly stirred. Mix and add another third of water and continue to cook and stir for about 45 to 60 minutes until a clear, orange-colored solution is obtained. Then add the rest of the water and the calcium arsenate. Let the mixture settle and run it through a fine sieve as it is poured into the spray tank. This should be diluted in a ratio of about six parts water to one part of the solution.

Soil Sterilization in Field and Garden Formula No. 1

The stand of such vegetables as peas, spinach and beets can usually be greatly improved by watering, immediately after planting, with a dilute solution of formaldehyde.

Formaldehyde (40%) 1 oz. Water 124 oz.

Use this solution at the rate of 1 gal. for 200 feet of row.

No. 2 Formaldehyde (40%) 15 oz. Infusorial Earth 85 oz.

When infusorial earth is used as a carrier the full strength of the formal-dehyde is maintained for a longer time than when other materials, such as charcoal or muck, are employed. Mix thoroughly, taking care to break up lumps. Use 6 oz. of this dust for each bushel of soil, or 1½ oz. per square foot of flat area. Insure that the dust is well mixed with the soil. After placing in flats, sow seed and water immediately.

Adhesives for Sulphur Dusts

Sulphur is more than twice as adhesive if applied to wet citrus foliage as if applied to dry foliage; 0.25 inch of rain removed so much sulphur dust applied to dry foliage that its effectiveness was lost. Addition of 2% of glue or gum tragacanth to dusting sulphur increased its adhesiveness 4-5 times over sulphur applied to dry foliage and twice over sulphur applied to wet foliage. When 5% of blood albumin was added to sulphur dust, its adhesiveness was increased 10 times and 5 times over that of sulphur applied alone to dry leaves and wet leaves respectively. Sulphur dust containing blood albumin remained on the leaves almost as well as did lime.

Pepper Disease Control

The use of an organic mercury dust or solution of 1 to 1000 mercuric chloride with an exposure of 5-8 minutes effectively sterilizes pepper seeds before planting. For treatment of the growing plant to control fungus diseases the use of either Bordeaux mixture or copper-lime dust is recommended. For the Bordeaux mixture, a concentration of 2-4-50 should be applied to seedbeds and 4-6-50 to more mature plants. The copper-lime dust should be mixed in the proportion of 20 lb. of dehydrated copper sulphate and 80 lb. of calcium hydroxide. These components should be mixed dry.

Dust for Control of Cucurbit Wilt

Basic Copper Chloride ½ oz.

Flour 5 oz.
Calcium Arsenate 1 oz.

Keep plants well covered with a light coating of dust from the time they appear through the ground until bearing stage is reached. New growth should be kept dusted. Number of applications will depend upon rate of growth and weather conditions.

Seed Disinfectant (Dustless) French Patent 770,560

Inloride	5	oz.
	5	oz.
	90	oz.
	Chloride	5

Lettuce Seed Sterilization
Soak 4 to 8 hours in following:
Calcium Hypochlorite 11.5 oz.
Water 1 gal.

Stir thoroughly; allow to settle; decant and use at once. Wash seeds after above treatment.

Spreader for Nicotine Sprays

Spreaders which contain twice the amount of active ingredients and which are 4 times as effective as potassium soaps in the control of Aphis Rumicis on nasturtium leaves, are made as follows:

Formula No. 1

Water 5 g., potassium hydroxide (92%) 7.40 g., pine-tar oil (specific gravity 1.035) 44.30 g., ethylene glycol monoethyl ether 10.00 g., oleic acid 33.30 g.

No. 2

Water 5 g., potassium hydroxide 7.40 g., pine-tar oil 48.80 g., isoamyl alcohol 3.00 g., phenol (85%) 1.00 g., ethylene glycol monoethyl ether 1.50 g., and oleic acid 33.30 g.

Cotton Root Rot Remedy Apply 3% ammonium hydroxide solution.

Preventing Brown-Rot on Lemon Trees
Apply

Zinc Sulphate 40–25 lb. Hydrated Lime 20–25 lb. Sand 40–50 lb.

around base of trunk, piling 8 inches high and hold in place by paper collar.

Lemon Scale Control

Yellow Sulphur		75 lb.
Gas Purification	Sulphur	25 lb.
Talc	-	10 lb.

Grind to 200-300 mesh; use 0.4 to 1 lb. per tree at 17-day intervals. Five to seven applications are used.

Control of Cabbage Root Fly

Corrosive sublimate, applied at a strength of 1 oz. in 8 gal. of water, is the most successful means, at present known, of reducing the damage done to plants of the cabbage tribe (Brassicæ) by the cabbage root fly. The treatment consists of applying to each plant about one-quarter of a pint of the solution in such a manner as to flood the soil evenly round the base of the plants on three occasions at 10-day intervals, starting four days after setting out the plants. Of the other methods tested, commercial naphthalene powder, about 1/4 oz., applied to the soil round the plants on three occasions at 10-day intervals commencing on the day of transplanting, possesses certain advantages, especially as regards cheapness, simplicity of application and the non-poisonous nature of the sub-

Control of Weevils in Stored Beans and Cowpeas

Protection is obtained by adding 1 lb. of slaked lime per bu. of beans or cowpeas and mixing thoroughly. Sodium fluosilicate, used at the rate of 1 part to 1500 parts of grain, gives full protection against the grain beetle, Sitophilus granaria.

Non-Poisonous Insect Exterminator

A TOTAL A GIROTAGUE ATTRECES	*****
Petrolatum, Liquid	1000 g.
Pyrethrum Powder	200 g.
Pine Needle Oil	13 g.
Juniper Oil	2 g.
Lavender Oil	1 g.
Orange Oil	1 g.

To Kill Ants in Lawns and Gardens

Make a hole about 18 inches deep in the center of the ant hill with an old broom handle and then pour in a solution of poison made by dissolving 1 oz. of sodium or potassium evanide * in 1 gal. of water. Cover with dirt. If the soil is alkaline use one-half the quantity of water and make another solution of 1 oz. of alum to 2 qt. of water and pour one-half of each in the hole.

* Deadly poison. Do not allow contact with broken skin or cuts.

Beetle Powders		Endive Fly Treatment
Formula No. 1		Soak roots, before planting, for 15 to
		20 minutes in:
Barium Carbonate	10 oz.	
Borax	20 oz.	Nicotine (50% Solution) 20 cc.
Sugar	5 oz.	Sal Soda 5 g.
No. 2		Water 1 l.
Sodium Fluoride	10 oz.	
Kaolin	10 oz.	Fly Dishes
No. 3		a. Quassia Wood 500 g.
Kieselguhr	22 oz.	Black Pepper 50 g.
Sodium Fluoride	40 oz.	Water 2 l.
Sodium Chloride	10 oz.	
		Extract cold 4 days, then evaporate to
No. 4		1 liter, filter and add:
Powdered Borax	4 oz.	b. Sugar 100 g.
Flour	2 oz.	Color with red or green aniline dye.
Chocolate Powder	1 oz.	Impregnate cardboard dishes with
No 5	0.0	solution; dry in air.
No. 5		boldston, dry in all.
Powdered Borax	10 oz.	Berging and Artifacture and Ar
Insect Powder	1 oz.	Killing Fly Larvae in Cesspools
Starch	1 oz.	Add 0.15% by weight of sodium cya-
And the second section of the s		nide to the fecal matter.
Poultry Lice Powde	are	
	CIS	70.
Formula No. 1		Derris Insecticide for Caraway Moth
Naphthalene	10 g.	Derris Root Powder
Sulphur	20 g.	(8% Rotenone) 1 kg.
Tobacco	40 g.	Tale 3 kg.
Talc	130 g.	Apply at rate of 75 kg. per hectare in
No. 2	· ·	two applications.
Naphthalene	20 g.	two approations.
Sulphur	20 g.	Section 19 Commence of the Com
Tobacco	40 g.	Grasshopper Poison
Talc	120 g.	Formula No. 1
No. 3		
Naphthalene	40 g.	
Sulphur	20 g.	Beet Molasses 2 gal.
Tobacco	40 g.	Amyl Acetate 3 oz.
Talc	100 g.	Sodium Arsenite, Liquid 1 qt.
No. 4		Water 10-12 gal.
Naphthalene	20 g.	No. 2
Sulphur	20 g.	Bran 100 lb.
Tobacco	40 g.	Sodium Arsenite, Liquid 1 qt.
Cresol	1 g.	Sodium Arsenite, Powder 2 lb.
Talc	119 g.	Water 10-12 gal.
No. 5	8.	part of the designation of the state of the
Naphthalene	20 g.	Groundnut (Peanut) Oil Insecticide
Sulphur	20 g.	
Sodium Fluoride	50 g.	This emulsion is made by mixing 500 cc. of groundnut oil with 75 cc. of oleic
Tale	110 g.	
	440 8.	acid and then pouring the mixture slowly,
Deer and Deulter Miss and	T in 17:11	with constant agitation, into a solution of
Dog and Poultry Flea and	Lice Killer	50 cc. of ammonia in 200 cc. of water.
Formula No. 1		For use, this emulsion should be diluted
Derris Powder	¼-1 kg.	with nine times its volume of water. It
Water	10 1.	is stated that all insects, the wax-covered
Shake well and rub into sl		bodies of which are resistant to ordinary
Shake well and lub into si	CIII.	aqueous liquids, are poisoned by this 2%
No. 2		oil emulsion.
Derris Powder	½-1 kg.	
Tale	10 kg.	Rat Fumigant
No. 3	~~g.	Potassium Nitrate 30 oz.
Rotenone Solution	0.2%	Sulphur 42 oz.
TACAGINATIO MAINTAIT	VIA 70	TA OLI

Sawdust Sand	18 6	
Mix together and burn.		
Rat Bait		
Formula No. 1		
Ground Dried Bread	65	lb.
Ground Fresh Pork Fat		lb.
Ground Fresh Halibut or Haddock or Cod		lb.
Powdered Red Squill	10	lb.
No. 2		
Ground Dried Bread	85	lb.
Glycerin		lb.
Powdered Red Squill	10	lb.
No. 3		
Ground Dried Bread	37	lb.
Glycerin		lb.
Powdered Red Squill		lb.
*Fresh Bait	50	lb.
No. 4		
Ground Dried Bread 10 lb.		
Corn Oil 1 lb. Zinc Phosphide	4 10	OZ.
	10	02.
No. 5		
Ground Dried Bread 5 lb.	10	
Corn Oil Zinc Phosphide	10 10	
Some Fresh Bait 6 lb.	4	OZ.
*		023
No. 6		
Ground Dried Bread 29 lb. Ground Fresh Pork Fat 2 lb.	0	oz.
Ground Fresh Halibut 6 lb.		
Powdered Thallium		
Sulphate	10	OZ.
No. 7		
Ground Dried Bread 18 lb.	2	oz.
Glycerin 1 lb.	4	oz.
Powdered Thallium	7.0	
Sulphate	10	
* Fresh bait has hamburger, or gro	und	sweet

* Fresh bait has hamburger, or ground sweet potatoes (raw but canned is better), or ground applies, or ground bananas.

Rat Poison for Flour Mills Sodium Silicofluoride 70 lb. Diatomaceous Earth 30 lb.

Dust on floor, keeping away from sacks. Rats lick powder off feet and go out seeking water and thus die outside.

Rabbit Poisons

Poisoned Alfalfa. Dissolve 1 oz. of strychnine sulphate in 1 gal. of hot water and sprinkle over 10 lb. of dry alfalfa leaves. Well-formed leaves free from dust or sticks should be used. They should be threshed thoroughly until all the moisture is absorbed. The poisoned leaves should be distributed in small handfuls, in lines a few feet apart, across portions of the field where observations show the rabbits to be feeding. Stock should be excluded.

Poisoned Green Alfalfa (summer poison)
Chopped Green Alfalfa 20 lb.
Strychnine (Powdered
Alkaloid) 1 oz.
Saccharine $\frac{1}{10}$ oz.

Poisoned Rye Heads. In localities where alfala is not raised, rye, emmer, or wheat heads are excellent mediums for poison, and frequently surpass alfalfa leaves in effectiveness, particularly in dry-land sections. Where possible, grain heads for poisoning should be cut and cured when the grain is in the dough stage, as it is more palatable and attractive to rabbits when cut at this time. Dissolve 1 oz. of strychnine sulphate in 6 qt. of hot water and sprinkle over 10 lb. of grain heads. Mix thoroughly until all moisture is absorbed. The heads should be cut from the stem just below the last kernel and as little straw taken as possible.

Cedar Shingles.

Strychnine (Powdered
Alkaloid)
Saccharine
Bicarbonate of Soda
(Baking Soda)
Flour

1 oz.

7 oz.

1 oz.

1 oz.

Mix together dry, 1 oz. of powdered strychnine (alkaloid), 1 oz. of baking soda, 1 teaspoonful of saccharine, and 3 tablespoonfuls of flour. Add a little cold water and stir thoroughly to a smooth, creamy paste. Split the shingles and dip the tops in the paste and stick them into the ground along the rabbit trails and runways. These baits can be easily taken up when they are no longer needed and all danger to stock is thereby eliminated. In many communities this poison has proved very effective.

Starch Formula (Rabbits). Dissolve 2 oz. (heaping tablespoonful) of gloss starch in a little cold water, pour into 2 to 3 quarts of boiling water, and stir inti a thin starch paste is formed. Stir into the starch paste 1 oz. of strychnine (alkaloid) until a creamy paste, free from lumps, is formed. Mix the paste thoroughly over 10 lb. of grain heads until every head is coated. The heads should be cut from the stem just below

the last kernel and as little straw taken as possible. Ten pounds of alfalfa leaves or chopped alfalfa may be used in place of grain heads in alfalfa districts.

Rabbit Salt. Mix dry 1 oz. strychnine (alkaloid) with 16 oz. granulated salt. A very satisfactory method is to bore about 2% of the way through a short 2" by 4" block with 1- to 1½-inch bit and place the salt bait in this container. The blocks should be placed in or near the rabbit trails and runways. Care should be taken in placing these baits so that livestock will not obtain them.

Insect Control in Stored Rice

Fumigate with 2.5 lb. chloropicrin per 1000 cubic feet of space at temperatures above 70° F. or with carbon bisulphide at rate of 6 lb. per 1000 cubic feet.

Ground Squirrel Poison

Mix thoroughly 1 oz. strychnine alkaloid (powdered) and 1 oz. baking soda. Sift this into ¾ pint of thin, hot starch paste and stir to a creamy mass. The starch paste is made by dissolving one heaping tablespoonful of dry gloss starch in a little cold water, which is then added to ¾ pint of boiling water. Boil and stir constantly until a clear thin

paste is formed.

Add ¼ pint heavy corn syrup and a tablespoonful of glycerin and stir thoroughly.

Add 1/8 oz. saccharine and stir thoroughly.

Pour this poison solution over 20 quarts of clean oats and mix thoroughly so that each grain of oats is coated. Prepare it 24 to 48 hours before using.

For mixing small quantities an ordinary galvanized wash tub is convenient. For large quantities a tight, smooth box may be used, and mixing may be done with a spade.

A teaspoonful of the poisoned oats should be placed near each ground squirrel hole on clean hard ground letting it scatter slightly as it falls. (Placed in this way it will not endanger stock). Do not put the poisoned grain on the loose dirt of the mound or into the holes. Each quart of the poisoned grain is sufficient to treat about 60 holes.

Squill Paste Preservative

A suitable preservative for the red squill paste is 1% of a hydroxybenzoic acid derivative or ½% of benzoic acid.

White Coal Tar Disinfectant

Cresylic Acid	50 lb.
Cresylic Creosote	6 lb.
Sulphonated Castor Oil	5 lb.
Gelatin	3 lb.
Water	36 lb.

The sulphonated oil and gelatin are dissolved in the water and the mixed tar acids are gradually added to them with vigorous agitation in small quantities at a time. Final treatment with a colloid mill may be necessary to obtain a good dispersion.

Insecticides to Be Applied by Fumigation

German Patent 597,769

Naphthalene

Formula No. 1

40 m

rapinnaiene	7:0	ц.
Naphthalene, Crude	40	g.
Animal Oil	5	cc
Cresol, Crude	5	cc
Ceresin	10	
No. 2		ρ,
110. 4		
Naphthalene	15	g.
Naphthalene, Crude	25	
Animal Oil	20	60
Cresol, Crude	30	
Ceresin		
	10	g.
No. 3		
Naphthalene	10	o٠.
Naphthalene, Crude		g.
Animal Oil	25	
Cresol, Crude	25	cc
Ceresin	30	ø.

Method: Melt naphthalene and ceresin, and add cresol and oil at low temperature. Application on fumigation-pans.

Warehouse Fumigant

Chloropicrin (nitrochloroform, CCl₃-NO₂) is a colorless heavy liquid which is becoming prominent as a fumigant. Due to its highly lachrymatory nature as well as its highly toxic effect on insects, their larvae and eggs, it makes possible effective fumigation without the high possibility of accidental death to operators attendant on the use of hydrogen cyanide.

The following are the recommended quantities in pounds per 1000 cu. ft. of volume to be fumigated using 24-hour exposure and a temperature of 70-80° F. Higher temperatures reduce the exposure time while lower ones increase it. The liquid is vaporized by spraying or evaporation from very shallow pans or soaked

cloth.

Confectionery Industry. Lb. per 1000 cu. ft.	{ Calcium Hypophosphite Sodium Hypophosphite Water, Distillate	8 g. 12 g. 71 g.
Candy 1 Nuts 1	No. 3	
Dairy Industry.	35% Oil	
Eggs and Cheese 14	Carragheen Moss	18 g.
Milling Industry.	Distilled Water	400 g.
	Sodium Formate Cod Liver Oil	5 g.
Macaroni Vaults 1-14 Macaroni, Cased 14-2	Syrup, White	350 g. 100 g.
Space Fumigation 1	Distilled Water	117 g.
Flour Mills	*Spice Oil Mixture	10 g.
General 1	•	8-
Returned Bags 11/4	No. 4	
Sacked Flour 11/2	30% Oil	19 ~
Fly and worm control.	Gum Arabic Gum Tragacanth	12 g. 16 g.
(Exposure over the week end) 1/4	Glycerin (28° Bé.)	140 g.
Rice Bags and Vaults 11/2	Distilled Water	430 g.
Box Cars (Adults) 4-5	Moldex or Other Good Pre-	•
Box Cars Complete 7-8	servative	1 g.
$Stored\ Products.$	Cod Liver Oil *Spice Oil Mixture	300 g.
Warehouse Space 1	Calcium Hypophosphite	10 g. 8 g.
Sacked Goods and Vaults 1½	Coding III	12 g.
Grain Bins	Distilled Water	71 g.
(With grain moving at 100	No. 5	
bu. per hr.) 2	Gum Arabic	15 g.
Contaminated Bins 3	Gum Tragacanth	8 g.
Tobacco.	Glycerin (28° Bé.)	50 g.
Vaults 2	Distilled Water	456 g.
Warehouses 11/4	Sodium Formate Iodine	5 g.
Furniture.	Chloroform	3 g. 3 g.
Furniture 11/4	*Spice Oil Mixture	3 g.
Household.	Cod Liver Oil	447 g.
Bedbugs, Clothes Moths,	No. 6	Ū
Roaches 1	Gum Arabic	10 g.
Buffalo Moth 11/4 Rodents 1/	Gum Tragacanth	10 g.
/±	Glycerin (28° Bé.)	200 g.
Cod Liver Emulsion for Animals	Water, Distilled Potassium Iodide	366 g. 3 g.
Formula No. 1	Moldex or Other Good Pre-	o g.
50% Oil	servative	1 g.
Carragheen Moss 12 g.	Cod Liver Oil	400 g.
Distilled Water 300 g. Moldex or Other Good Pre-	*Spice Oil Mixture	10 g.
	No. 7	
servative 1 g. Cod Liver Oil 500 g.	Carragheen Moss	19 g.
Syrup, White 86 g.	Glycerin (28° Bé.)	100 g.
Distilled Water 91 g.	Distilled Water	519 g.
*Spice Oil Mixture 10 g.	Potassium Iodide Moldex or Other Good Pre-	1 g.
No. 2	servative	1 g.
40% Oil	Cod Liver Oil	350 g.
Gum Arabic 12 g.	*Spice Oil Mixture	10 g.
dum Tragacantn 12 g.	No. 8	-
Glycerin (28° Be.) 130 g.	Gum Arabic	12 g.
Water, Distilled 340 g. Sodium Salicylate 5 g.	Gum Tragacanth	16 g.
Sodium Salicylate 5 g. Cod Liver Oil 400 g.	Glycerin (28° Bé.)	130 g.
*Spice Oil Mixture 10 g.	Distilled Water Sodium Salicylate	426 g.
* ** ** ** ** ** ** ** ** ** ** ** ** *	1 Southin Sancylate	5 g.

Iodine	1 g.
Alcohol, Absolute	10 g.
Cod Liver Oil	300 g.
Spice Oil Mixture	10 g.
Calcium Hypophosphite	8 g.
Sodium Hypophosphite	12 g.
Calcium Hypophosphite Sodium Hypophosphite Distilled Water	70 g.
*Snice Oil Mixtures for Ahove	Emulsion

Spice Oil Mixtures for Above Emulsions

Formula No. 1

Vermouth Oil	5	cc.
Coriander Oil	2	cc.
Galanga Oil	1	cc.
Gentian Oil	1	cc.
Calamus Oil	0.5	
Peppermint Oil	0.5	cc.

No. 2

5 (зc.
3 (зc.
2 (e.
	3 (

No. 3

Fennel Oil	6	cc.
Calamus Oil	4	cc.

The above emulsions are made up best in enameled kettles with high speed mixers.

Gum Solutions: Wash gum arabic with water at 40° C., then put into cold water and warm to solution. Gum tragacanth or carragheen moss are first wet with glycerin and put into cold water. Soak 12 hours. Prepare gums separately and when ready, mix as indicated, warm up to 90° C. (add Iodide) then add preservative.

Stir in Cod Liver Oil in small portions. Then add Spice Oil Mixture, with stirring. Syrup and Hypophosphites are dissolved in hot water as indicated. Stir into emulsion hot. Iodine is prepared by solution in Alcohol or Chloroform and a little of the Cod Liver Oil, then is added to the gum (aqueous) solutions and emulsified.

When ready, stir vigorously for 1/2 hour, or put through a homogenizer.

Cod Liver Oil Emulsion for Animals

Formula No. 1

Gum Arabic	100	g.
a. Gum Tragacanth	100-120	
Glycerin	1200	g.
b. Cod Liver Oil, Crude	3700	g,
Calcium Hypophosphite	50	g.
c. Sodium Hypophosphite	50	
Water	4000	ø.

Grind a until smooth, add b in small portions, homogenizing every time. To this add c in an emulsifying machine.

As spice, add 1% of the following mixture of:

Vermouth Oil	10 cc.
Coriander Oil	4 cc.
Galanga Oil	2 cc.
Gentian Oil	2 cc.
Calamus Oil	1 cc.
Peppermint Oil	1 cc.

No. 2

[Iceland Moss		10	g.
a. { leeland Moss } Water (2 portions)	to	600	g.
			Extract
$b. egin{cases} \operatorname{Gum} & \operatorname{Tragacanth} \\ \operatorname{Gum} & \operatorname{Arabic} \\ \operatorname{Cod} & \operatorname{Liver} & \operatorname{Oil} \end{cases}$		6	g.
b. Gum Arabic		6	g.
Cod Liver Oil		400	g.
c. Fennel Oil		5	g. drops
Calamus Oil		5	drops

Boil a two times (two portions of water) to 600 g. united extract. Grind b until homogeneous and transfer into a dry bottle; add c, then a in two portions, shaking thoroughly and vigorously.

No. 3

a. Carragheen Moss	10 g.
Water	350 g.
b. Cod Liver Oil	500 g.
$c. \begin{cases} \text{White Syrup} \\ \text{Malt Extract} \\ \text{Water} \end{cases}$	100 g.
c. { Malt Extract	29 g.
Water	120 g.

Soak a for 12 hours, boil then about 10-15 min., filter through cloth. Add b, while stirring, to this hot solution, then stir in c, and add as preservative

Sodium Salicylate 0.3-0.5 g.

No. 4

a.	Gum Tragacanth	5	g.
	Gum Arabic	8	g.
	Water	250	g.
Ъ.	Calcium Chloride	50	g.
	Water	57	ğ.
c.	Lime Water	230	g.
d.	Cod Liver Oil	400	g.

Soak a for $1\frac{1}{2}-2$ days, add b, then c, mix well, percolate (lumps remaining on the cloth are ground with water and pour again through the filter). Mix the whole well in an emulsifying machine with d for hours; d is added in 8 portions. The a, b, c is treated alone before.

Skin Abrasion Lotion (For Dogs)

Dissolve 1 part of castile soap in 9 parts of water. Wash dog thoroughly with this solution; and then apply with cotton to the affected parts 5% tincture of iodine.

Moisture Eczema Lotion for Dogs
This lotion is excellent for bathing

moist eczema spots on dogs.

_	0	
Tannic Acid	5	oz.
Salicylic Acid	5	OZ.
Alcohol (50%)	90	oz.

Before using this preparation the spots should be thoroughly washed with castile soap.

Dog Eczema Powder

Senega Root Powder	90 oz.
Sodium Sulphite	10 oz.

Rub into skin with water and finally wash off.

Liquid Soap for Dogs and Other Animals

Olein Caustic Potash (50%) about 736 g. Glycerin Softened or Distilled Water 6800 g. Carbolic Acid (Phenol),	Palm Kernel Oil	1200	g
Glycerin 600 g. Softened or Distilled Water 6800 g.	Olein	300	g.
Softened or Distilled Water 6800 g.	Caustic Potash (50%) abou	t 736	g.
Softened or Distilled Water 6800 g.	Glycerin	600	g.
Carbolic Acid (Phenol),	Softened or Distilled Water		
			3

Crude 400 g.
Perfume Oil (e.g., Eucalyptus) 50 g.

Dog Deterrent

Naphthalene Flakes	4 oz.
Paraffin Wax	1/4 oz.
Gasoline	1-2 pt.
Rosin	1/4 07

Stir until dissolved; spray on base of tree trunks or shrubs with an insect spray gun.

Dog Nuisance Preventer

To prevent dogs from staining trees and shrubs, spray the base of the latter with a solution of ¼ oz. nicotine sulphate per gal. water.

Dog Worm Remedy Formula No. 1

Aloes	45 gr.
Soap	45 gr.
Oleoresin of Male Fern	30 gr.
Mix and make into 2 pills.	

Administer both pills in the morning, the animal to remain fasting for some time.

No. 2

Areca nut, freshly ground, is considered an excellent remedy for worms in dogs. About one dram made into a pill is the dose for an ordinary sized dog. This should be given at night followed by a dose of castor oil in the morning.

Animal Eye Washes

One of the best eye washes for irrigation and cleansing of the eye and for purulent discharges and conjunctivitis is as follows:

Sodium Bicarbonate	15 gr.
Borax	15 gr.
Sodium Chloride	15 gr.
Glycerin	1 dr.
Distilled Water	8 oz.

Animal Ear Preparation Formula No. 1

Gentian	Violet	5	oz.
Acetone		5	oz.
Alcohol		45	oz.
Water		45	oz.

Take small amount in an eye dropper and place deep into the ear and remove excess so as not to soil the outside.

No. 2

Phenol	3	OZ.
Glycerin	97	OZ.

Add boric acid powder until the glycerin will not absorb any more. Let stand over night and strain.

Place one-half eye dropperful in ear and remove the excess.

Dog Mange Treatment

Formula No. 1

Kerosene	32	oz.
Creolin	6	oz.
Oil of Tar	6	oz.
Sulphur	1	lb.
Raw Linseed Oil	to make 1	gal.

Rub into skin every other day. It gives gratifying results.

No. 2

Another good oily skin mixture is:

Gum Camphor	1 lb.
Alcohol	1 pt.
Turpentine	1 qt.
Kerosene	2 qt.
Cotton Seed Oil	6 qt.
Sulphur (Flowers)	9 oz.

Note: First dissolve the camphor in the alcohol. Rub on the skin every third day.

Dog Mouth Wash

Tincture Iron			1	OZ.	
Potassium Chlorate			2	oz.	
Glycerin			4	OZ.	
Water	to	make	1	gal.	

Aphrodisiac for Cattle and Horses

The usual doses of yohimbine hydrochloride as an aphrodisiac in veterinary practice are: Stallions, 1 gr.; bulls, 1¼ gr.; cows and mares, 1½ gr. It should be administered in the food or drinking water three times a day.

Cow Abortion Flush

Common Salt	1	lb.
Potable Water	. 95	lb.

Remove aborting cow from herd. Before returning to herd flush daily with above solution.

Bloody Milk Mixture

Glauber's	Salts	1	lb.
Water		4	lb.

Give the above dosage to cow producing bloody milk. Find and remove the cause; it may be udder injury, improper feeding, or overfeeding. Certain bacteria impart a red color to milk; this is uncommon.

Cow Boil Wash

Carbolic Acid Solution (3%)

Syringe out cavity with above solution after lancing and removing contents.

Chapped Teats Solution

Boric Acid	Crystals	1	lb.
Water		15	lb.

Bathe teats twice daily with above and dry; then rub teats with vaseline.

Cow Pox Solution

Apply a 4% solution of potassium permanganate after cleaning udder and teats.

Calf Scours Remedy

Salol			1	lb.
Subnitrate	of	Bismuth	2	lb.

First give the calf with simple scours 1½ oz. of castor oil in ½ pt. of warm milk. After a few hours give a teaspoonful of the above. Repeat this dosage three times daily.

Impaction in Cattle Treatment

Glauber's Salts	11/2	lb.
Water	7	lb.

Administer 2 oz. of aromatic spirits of ammonia at once. Two hours later give the above formula.

Egg Preserving Solution

Sodium	Silicate	1	fl.	oz.
Water		25	fl.	oz.

Defeathering Poultry U. S. Patent 2,017,648

Burgundy Pitch	15 lb.
Montan Wax	5 lb.
Paraffin Wax	10 lb.

The ingredients are melted, thoroughly mixed, and applied to the carcass, preferably after the bird has been scalded and the bulk of the feathers that can be removed hastily have been removed by hand.

After application, the defeathering compound is permitted to solidify by cooling and is then removed, taking with it epidermal excrescences such as feathers, down, pinfeathers and the like.

Prevention of Skin Tearing when Plucking Feathers with Adhesives

Spray the skin with an oil emulsion.

Bird Gravel Fine River Sand 97.5 g. Cuttlefish Bone, Powder Pyrethrum Flowers 2 g. 0.5 g.

Ration No. 1 (With Milk)

For those who wish to use home-grown grains.

grams.		
Ground Corn	18	lb.
Ground Barley	18	lb.
Ground Wheat	18	lb.
Ground Oats	18	lb.
Meat Scrap	10	lb.
Dried Milk	10	lb.
Alfalfa Meal	5	lb.
Steamed Bone Meal	2	lb.
Salt	1	lb.
T		

Ration No. 2 (With Milk)

Using wheat by-products.	
Ground Corn	20 lb.
Bran	20 lb.
Flour Middlings	19 11

 Ground Oats
 20 lb.

 Meat Scrap
 10 lb.

 Dried Milk
 10 lb.

 Alfalfa Meal
 5 lb.

 Steamed Bone Meal
 2 lb.

 Salt
 1 lb.

This is a ration for those who wish to use barley in the laying ration. Barley

is not as palatable as corn when fed whole in the scratch grain but is a valuable ingredient of a laying mash. However, it should be remembered that this grain is low in vitamin "A" when compared with corn and that sufficient alfalfa meal should be present to take care of this deficiency.

Ration No. 3 (With Milk)

(
Ground Barley	20 lb.
Bran	20 lb.
Flour Middlings	10 lb.
Ground Oats	20 lb.
Meat Scrap	10 lb.
Dried Milk	10 lb.
Alfalfa Meal	7 lb.
Steamed Bone Meal	2 lb.
Salt	1 lb.

Ration No. 4 (Without Milk)

(
Ground Corn	20	lb.
Bran	20	lb.
Flour Middlings	14	lb.
Ground Oats	20	lb.
Meat Scrap	20	lb.
Alfalfa Meal	5	lb.
Salt	1	lb.

Ration No. 5

Many farmers and poultrymen wish to feed a surplus of liquid milk (either skim or buttermilk) to the laying flock. This is a successful practice and the following ration is designed to be fed when liquid milk is given as the only drink. omitting the water for drinking purposes no fear need be felt as liquid milk is about 90% water. If water is given to the flock in addition to the liquid milk, the meat scrap content should be increased to obtain best results. It should also be remembered that the practice of feeding liquid milk for one or two days and then missing a day is a bad one and a satisfactory production cannot be expected over a long period of time.

Ground Corn		20	lb.
Ground Oats		21	lb.
Ground Barley		21	lb.
Ground Wheat		20	lb.
Meat Scrap		10	lb.
Alfalfa Meal		5	lb.
Steamed Bone	Meal		lb.
Salt		1	lb.

The above ration may also be used when condensed buttermilk is fed.

Ration No. 6

The following ration is one which has been fed at the Poultry Experiment Station to 1200 laying hens during the past

year and the egg production and hatchability obtained have been satisfactory.

Ground Barley	28 lb.
Bran	20 lb.
Ground Oatmeal	11 lb.
Flour Middlings	10 lb.
Meat Scrap	10 lb.
Dried Milk	10 lb.
Alfalfa Leaf Meal	8 lb.
Steamed Bone Meal	2 lb.
Salt	1 lb.

Either dried buttermilk or dried skim milk may be used in making up these laying rations. If one is feeding for eggs alone, any of the foregoing rations will give good results. If good hatchability is desired, rations No. 1, No. 2, No. 3 and No. 6 are recommended.

A satisfactory scratch grain consists of equal parts, by weight, of corn and wheat.

It is important that pullets especially obtain enough scratch grain to keep them in good growing condition. They are under the double strain of egg production and growth. Do not obtain fall and winter eggs from your pullets at the expense of growth as this leads to moult.

Oyster shell should be available to the laying flock at all times.

All-Mash Ration for Laying Hens Yellow Corn, Coarsely Ground lb. Wheat, Coarsely Ground or Shorts 30 lb. Oats, Finely Ground 20 lb. Wheat Bran, Coarse lb. Meat Scrap, Medium (50-55% Protein) 10 lb. Dried Skim Milk or Buttermilk lb.

In addition, for confined layers, use ½ to 1 pt., or amount suggested by manufacturers, of potent cod liver oil or sardine oil to each 100 pounds of mash.

5 lb.

Alfalfa Meal or Leaf Meal

Salt

In case it is preferred that grain and mash be fed separately, the following formulas may be used:

Mash Ration

	Coarsely Ground Yellow Co	rn	20	lb.
7	Vheat, Coarsely Ground or			
	Shorts		20	lb.
C	Oats, Finely Ground		20	lb.
7	Vheat Bran, Coarse		9	lb.
Ŋ	Meat Scrap, Medium			
_	(50-55% Protein)		20	lb.
·I	Oried Skim Milk or Butter-			
	milk		- 5	lb.

Alfalfa Meal Salt	5 lb. 1 lb.
Cod Liver Oil	1 lb.
Grain Ration	
Whole Wheat Whole or Cracked Corn Whole Oats or Barley	2 lb. 2 lb. 1 lb.
Whole Wheat Whole or Cracked Corn }	equal parts

Egg-Laying Rations Mixture No. 1

Mash:		
Corn Meal		16 lb.
Meat Scrap		6 lb.
Bran		1 lb.
Middlings		1 lb.
Scratch Mixture:		
Cracked Corn		1 lb.
Wheat		1 lb.
	No. 2	
Mash:		

Mash:	
Barley Meal	2 lb.
Bran	1 lb.
Middlings	1 lb.
Fish Scrap	1 lb.
Scratch Mixture:	
Cracked Corn	1 lb.
Wheat	1 lb.

With the above mixtures supply some green feed. Feed scratch mixture twice daily and sparingly. Feed scratch mixture early in the morning and late in the afternoon. Mash may be fed dry or wet.

Chick Feed

Yellow Corn Meal (Ground	
Coarsely)	360 lb.
Bran	200 lb.
Ground Oatmeal	200 lb.
Skim Milk Powder	100 lb.
Meat Scrap	50 lb.
Alfalfa Leaf Meal	50 lb.
Steamed Bonemeal	20 lb.
Salt	10 lb.
Cod Liver Oil	1 lb.

Chick Starter Feed

Yellow Corn, Coarsely Ground Coarsely Ground Wheat or	50	lb.
Middlings Wheat Bran		lb.
Meat Scrap (50-55% Pro-	-	
tein)	10	lb.

Dried Skim Milk or Butter-	
milk	5 lb.
Alfalfa Meal or Leaf Meal	5 lb.
Cod Liver Oil	1 lb.

Ration for Fattening Chickens Formula No. 1

Finely Ground Corn	12	lb.
Wheat Bran	4	lb.
Wheat Middlings	4	lb.
Meat Scrap	1	lb.
No. 2		

No. 2		
Finely Ground Oats	15	lb.
Finely Ground Corn	15	lb.
Low Grade Flour	2	lb.
Bran	1	lb.

To fatten chickens, feed one of the above mixtures 3 times daily. Food should be made soft with buttermilk or skim milk.

Breeding Flock Ration

Mash:		
Bran	1	lb.
Middlings	1	lb.
Corn Meal	3	lb.
Meat Scrap	11/2	lb.
Ground Oats	1	lb.
Rolled Oats	1	lb.
Linseed Meal	1/2	lb.
Scratch Mixture:		

Scratch Mixture:	
Cracked Corn	1 lb.
Wheat	1 lb.

Keep breeding stock outdoors every good day throughout the year. Supply abundance of green feed. Feed scratch feed in deep litter to make hens exercise. Fertile eggs can be produced by not forcing the hens with food, and by keeping vigorous males also well fed.

Poultry Appetite Stimulant

round Appende	Cumulant		
Pulverized Gentian	1	lb.	
Pulverized Ginger	1/4	lb.	
Pulverized Saltpeter	1/4	lb.	
Pulverized Iron Sulph	nate ½	lb.	
Pulverized Nux Vomi	ca 1/4	lb.	
4777 0 17			

Add 1 oz. of the preparation to each 5 lb. of mash.

Poultry Coccidiosis Feed

Dry Skim Milk or Butter-		
milk	40	lb.
Wheat Bran	10	lb.
Yellow Corn Meal	30	lb.
Ground Barley	20	lb.
Ferrous Sulphate	1/4	lb.

20 g.

Powder for Hens to Increase Egg Production

	~		
Formula	No. 1	No. 2	No.
Dicalcium Phosphate,	g.	g.	g.
Precipitated	72	70	
Calcium Carbonate			60
Ferrous Sulphate, Pow-			
der	12	10	
Ferrous Oxide, Powder.		-	10
Black Pepper, Ground	6	_	5
Ginger Root, Powder	_	20	10
Gentian Root, Powder	10		
Stinging Nettle Seed		-	15

Harrison Test Cow Feed

This is the formula recommended by Cornell University for test cows. It can be successfully used with second cutting alfalfa or second cutting timothy and clover.

Formula No. 1

Distillers Grain (9% Fat)	300 lb.
Wheat Bran	400 lb.
Hominy or Corn Meal	400 lb.
Ground Oats	370 lb.
Coconut Oil Meal	300 lb.
Linseed Oil Meal	200 lb.
Steam Bone Meal	20 lb.
Salt	10 lb.
(18% protein feed.)	

No. 2

Soybean Feed

This can be successfully fed with good hay.

Ground	Oats	900	lb.
Ground	Soybeans	100	lb.

Fattening Powder for Pigs

Formula	No.	No. 2	No.
	g.	g.	g.
SaltAntimony Sulphide	11	20	10
(Sb ₂ S ₃ crude)	10	10	-
Sulphur Flowers	11	10	_
lized	11	20	10
Sodium Bicarbonate	21		
Trigonella Seed	16	10	20
Linseed Meal	20	10	
Fennel, Pulverized		10	 13
Gentian Root Powder		10	13
Juniper Berries, Dry, Pow-		y - 1	
der	-	10	20
Calamus, Powder	-	-	14

Milk-Increasing Powder for Cows Formula No. 1

50 g.
30 g.
20 g.
40 g.
20 g.
20 g.

Goat Feeds

Trigonella Seed

1. Ground Feed for Bucks

Ground Corn	100 lb.
Ground Oats	100 lb.
Bran	50 lb.
Linseed Meal	25 lb.

Feed at rate of 1½ lb. per buck daily; increase to 2 lb. during breeding season. Include 3 lb. of alfalfa or clover hay, and a pound of turnips with the ration of ground feed.

2. Vorhies Grain Mixtures for Does

I.	
Rolled Barley	100 lb.
Wheat Bran	100 lb.
Dried Beet Pulp	100 lb.
Coconut Oil	100 lb.

Feed 1 to 2 lb. per doe daily along with hay and mangels.

II.

Dried Beet Pulp	600 lb.
Rolled Barley	100 lb.
Wheat Bran	100 lb.
Coconut Oil	200 lb.

Feed 1 to 2 lb. per doe daily along with hay and turnips.

111.	
Dried Beet Pulp	100 lb.
Wheat Bran	100 lb.
Oats	100 lb.
Coconut Meal	100 lb.

Feed 1 to 2 lb. per doe per day along with hay and turnips or silage.

Dried Beet Pulp	300 lb.
Rolled Barley	100 lb.
Wheat Bran	100 lb.

Feed 1 to 2 lb. per doe per day along with hay and mangels.

California Kid Feed Formula

Rolled Barley	100 lb.
Ground Oats	100 lb.

Feed 1/4 to 1/2 lb. daily per kid after two weeks of age. Allow animals to eat hay, and give milk.

Feeding Lime for Animals

Formula	No. 1	No. 2	No.
Dicalcium Phosphate, Pre-	g.	g.	g.
cipitated	65	70	80
Salt	10		5 3
Licorice Root, Powder	6	9	3
Calamus, Pulverized	4	4	3
Fennel, Ground Finely	4	4	3
Juniper Berries, Dry,		1	1
Powder	4	3	3
Trigonella Seed	7	9	3

Pasture Seed Mixture

Formula No. 1

Timothy	40 lb.
Alsike Clover	20 lb.
Kentucky Bluegrass	20 lb.
White Clover	20 lb.
Orchard Grass	20 lb.
Redtop	20 lb.
Meadow Fescue	20 lb.

The above formula is used for seeding pastures not to be hayed. Use 16 lb. of formula per acre.

No. 2

For Wet and Unproductive	Land
Alsike Clover	20 lb.
Canada Bluegrass	40 lb.
White Clover	20 lb.
Orchard Grass	40 lb.
Redtop	40 lb.
Use 16 lb. per acre.	

No. 3

110. 0	
Timothy	80 lb.
Red Clover	20 lb.
Alsike Clover	20 lb.
Kentucky Bluegrass	20 lb.
White Clover	20 lb.
Redtop	20 lb.
Orchard Grass	20 lb.

Use 20 lb. per acre. For a year or two the field should be hayed. After that when the plants are firmly established it should be pastured.

Garden Fertilizer

ATTLANTO DE CINTO	135 lb.
Nitrate of Soda	
Sulphate of Ammonia	200 lb.
Animal Tankage	250 lb.
Superphosphate	1000 lb.
Muriate of Potash	200 lb.
Filler	215 lb.

Fertilizer

Formula No. 1

French	Patent	779,281		
Calcium Phos	phate		75	lb.
Gypsum	~		20	lb.
Sulphur			5	lb.

No. 2

U. S. Patent 1,931,296

Roast following mixtures for 30 minutes at 315-425° F.

a. Rock Phosphate	40	lb.
Lime	10	lb.
Salt	$2\frac{1}{2}$	lb.
b. Coal	35	lb.
Salt	$2\frac{1}{2}$	lb.
Grind above with		
Ammonium Sulphate	10	lb.

No. 3

British Patent 410,487

Moist Sewage Sludge	50	lb.
Chalk	15	lb.
Slaked Lime	5	lb.
Dust, Refuse, Etc.	30	lb.

No. 4

U. S. Patent 2,019,713

Ammonium nitrate and ammoniated triple superplosphate in the proportions of about 45 to 60 parts of ammonium nitrate to 55 to 40 parts of ammoniated triple superplosphate.

Plant Food

Trisodium Phosphate	2	oz.
Potassium Sulphate	2	0 z.
Sodium Nitrate	3	oz.

Grind together and mix well. Only about a half gram of the above mixture should be used per plant every month or two. Caution: Using too much of any plant food is dangerous.

House Plant Food

Potassium Nitrate (Salt-

peter) 3 oz. Tribasic Sodium Phosphate 2 oz.

Mix, and dissolve about one tablespoon to the gallon. Of this solution, use one gill for each average size plant, once every two weeks.

Alkali Farm Land Treatment

"Alkali" spots on western farm land are usually due to the presence of sodium clay. Finely pulverized gypsum (calcium sulphate), thoroughly worked into the soil over a period of a year, will usually prove an effective remedy.

Detecting Treated Grains

Limed grain may be easily detected by the red color developed when it is dropped into a dilute solution of phenolphthalein.

Sulphur bleached grain may be detected by the dark color developed when

it is dropped into a dilute solution of lead acetate or lead nitrate.

Delinting Cotton Seed

Seed having a moisture content of 7 to 10% is treated with hydrochloric acid (2% on weight of seed) up to 60° E for 7 minutes. Treatment at 20° E requires 15 to 30 minutes.

FOOD PRODUCTS, BEVERAGES, FLAVORS

Ice Cream

Formulas are presented for seven series of ice-cream mixes containing 20 to 50% cream, showing the proportions of whole, skimmed, condensed, or dried milk that must be mixed in various combinations to produce the desired percentage of solids in the ice cream. These formulas show the ratios of milk fat to serum solids which are commonly used for different types of ice cream.

In Tables 1 to 5, the formulas contain the following dairy products, with 15% sugar added to the mixtures and 0.3% gelatin:

- Cream, skim milk, and whole milk.
 Cream, unsweetened condensed skim milk, and either skim milk or whole milk.
- (3) Cream, dry skim milk, and either skim milk or whole milk.
- (4) Cream, sweetened condensed whole milk and skim milk.
- (5) Cream mixed with 50% butter, to which mixture is added either dry or unsweetened condensed skim milk, making ice cream containing about 6 to 11% butter.

Tables 6 and 7 show combinations of dairy products without the addition of sugar which are suitable for basic mixes in milk plants for shipment to ice cream manufacturers. In each case, 15 lb. of sugar should be added to each 85 lb. of the unsweetened mix, in order to make a palatable commercial product. The dairy products used in these two tables are:

(6) Cream, unsweetened condensed skim milk, and either whole or skim milk.

(7) Cream, dried skim milk, and either whole or skim milk.

For each of these formulas there are given:

- (A) Percentage of solid constituents desired in the ice cream to produce ice cream containing 10 to 18% fat, from 20 to 50% ice cream;
- (B) Groups of ingredients which may be used in making comparable ice creams of the same solids content;
- (C) The percentages by weight of each of the different milk products required to give a mixture of the desired solids content.

The quantity of each ingredient needed for different size batches of the various mixtures can easily be determined by multiplying the quantity of the total mixture desired in pounds, by the percentages given in the tables.

The flavor and texture of ice cream will vary according to the proportion of milk solids, sugar, gelatin, and flavoring materials present, and the quality of the ingredients used. Ice-cream makers should therefore be careful to select ingredients and ice-cream formula of character and type best suited to their trade, and should check the accuracy of their figures in proportioning each mixture.

Ice Cream

Formula No. 1

100 lb. Mix—8% Fat—Cream, Whole Milk and Skim Milk Powder, 12%

~ugu		
Sugar	12	lb.
Gelatin	0.5	lb.
Skim Milk Powder	6	lb.
Cream (30%)	18.5	lb.
Milk (4%)	63	lb.
No 2		

8% Fat—Cream Skim Milk, Skim Milk, Powder, 12% Sugar.

Sugar	12	lb.
Gelatin	0.5	lb.
Skim Milk Powder	6	lb.
Cream (30%)	26.7	lb.
Skim Milk	55.8	lb.

No. 3

8% Fat—Sweet Butter, Whole Milk, Skim Milk Powder, 12% Sugar.

DAIM MILE I OFFICE, IM/O	Nugar	
Sugar	12	lb.
Gelatin	0.5	lb.
Skim Milk Powder	6	lb.
Butter (84%)	6	lb.
Milk (4%)	75.5	lb.
No 4		

8% Fat—Sweet Butter, Skim Milk, Skim Milk Powder, 12% Sugar.

Sugar	12	lb.
Gelatin	0.5	lb.
Skim Milk Powder	6	lb.
Butter (84%)	9.6	lb.
Skim Milk	72	lb.

-Amounts of Crosm of Different Fot Content and Fither Skim Milk or

J	ADIJE			Necessary									
		VV 11016	TATILL	recessary	101	making	Different	туј	pes ()I ICG	Oream		
,	g _o	lide					Tymog of	Tac	Cro	nam			

	a Per Cent	b Per Cent	c Per Cent	d Per Cent	e Per Cent	
Fat	14 6.39 15.00 0.3 35.69	16 6.30 15.00 0.3 36.60	16 6.21 15.00 0.3 37.51	$\begin{array}{c} 17 \\ 6.12 \\ 15.00 \\ 0.3 \\ 38.42 \end{array}$	18 6.03 15.00 0.3 39.33	
B. Ingredients Per cent (C. Percentage		dient by W Cream Mix		Each Type	of
	a	\boldsymbol{b}	\boldsymbol{c}	d	e	
1 Cream 50 Skim Milk 0	$28.00 \\ 57.00$	30.00 55.00	$\frac{32.00}{53.00}$	$34.00 \\ 51.00$	$\frac{36.00}{49.00}$	
2 Cream 40 Skim Milk 0	35.00 50.00	$37.50 \\ 47.50$	$\frac{40.00}{45.00}$	$42.50 \\ 42.50$	$\frac{45.00}{40.00}$	
3 Cream 30 Skim Milk 0	$\frac{46.75}{38.25}$	50.00 35.00	$53.50 \\ 31.50$	$56.75 \\ 28.25$	$\frac{60.00}{25.00}$	
4 Cream 20 Skim Milk 0	$70.00 \\ 15.00$	$75.00 \\ 10.00$	80.00 5.00	85.00		
5 Cream 50 Whole Milk 4	$23.00 \\ 62.00$	$25.25 \\ 59.75$	$27.50 \\ 57.50$	29.75 55.25	$\frac{32.00}{53.00}$	
6 Cream 40 Whole Milk 4	29.5 55.5	32.25 52.75	$35.00 \\ 50.00$	$37.75 \\ 47.25$	$\frac{40.50}{44.50}$	
7 Cream 30 Whole Milk 4	40.75 44.25	$\frac{44.75}{40.25}$	48.50 36.50	52.50 32.50	56.50 28.50	
8 Cream 20 Whole Milk 4	66.75 18.25	72.50 12.50	78.75 6.25	85.00	28.50	
Add to each above combination:			,			
Sugar	15.00 0.3	$\begin{array}{c} \textbf{15.00} \\ \textbf{0.3} \end{array}$	$\begin{array}{c} 15.00 \\ 0.2 \end{array}$	15.00 0.3	$\frac{15.00}{0.3}$	
Total	100.00	100.00	100.00	100.00	100.00	

Note: Ice creams made from these formulas whip and freeze slowly, and are likely to develop a buttery consistency, especially if the temperature is not kept fairly constant during storage in the hardening room or cabinet. The use of homogenized cream or mix will prevent undesirable fat clumping in freezing. Aging of the mixes for 24 hours at 40-50° F. before freezing will improve the texture. The cream flavor will be especially noticeable in the high-fat ice creams, hence care should be taken to use only high-grade cream. Melting will be accompanied by leaking of a milky serum from the ice and whipped cream structure of these ice creams, which keep their original form to a considerable extent instead of melting in a homogeneous mass. This is a natural characteristic of straight-cream ice creams, and does not constitute a defect.

No. 5		No. 6		
8% Fat—Sweet Butter, Skim Mil der, Water, 12% Sugar.	k Pow-	8% Fat—Cream, Whole Mill Powder, 14% Sug		Milk
Butter (84%) 9 Skim Milk Powder 13	lb. .5 lb. .6 lb. .2 lb. .7 lb.	Sugar Gelatin Skim Milk Powder Cream (25%) Milk (4%)	0.5 I 6 I	b. b.

Table 2.—Amounts of Cream of Different Fat Content; Unsweetened Condensed Skim Milk, and Fresh Skim Milk or Whole Milk Necessary for Making Different Types of Ice Cream

A. Solids		Types	of Ice Cr	eam	
	a	ъ	c	$ ilde{a}$	e
	Per Cent	Per Cent	$\operatorname{Per}\operatorname{Cent}$	Per Cent	Per Cent
Fat	10 11 15	$12 \\ 10 \\ 15$	14 9 15	16 8 15	18 7 15
Gelatin Total Solids	$\begin{array}{c} 0.3 \\ 36.3 \end{array}$	$\begin{matrix} 0.3 \\ 37.3 \end{matrix}$	$\begin{array}{c} 0.3 \\ 38.3 \end{array}$	$0.3 \\ 39.3$	$0.3 \\ 40.3$
B. Ingredients Per cent C. fat	Percentage		lient by W Cream Mix		Each Type of
	a	\boldsymbol{b} .	c	d	ϵ
1 Cream 50 Skim Milk 0	$20.00 \\ 41.00$	$24.00 \\ 42.50$	$\frac{28.00}{42.50}$	$\frac{32.00}{43.00}$	$36.00 \\ 43.75$
2 Cream 40 Skim Milk 0	$25.00 \\ 36.00$	30.00 36.50	35.00 35.50	$\frac{40.00}{35.00}$	$45.00 \\ 34.75$
3 Cream 30 Skim Milk 0	$33.30 \\ 27.70$	$\frac{40.00}{26.50}$	$\frac{46.70}{29.80}$	53.40 21.60	60.00 19.75
4 Cream 20 Skim Milk 0	50.00 11.00	$60.00 \\ 6.50$	$70.00 \\ 0.50$		
5 Cream 50 Whole Milk . 4	$16.5 \\ 44.5$	$20.50 \\ 46.00$	$\frac{24.50}{46.00}$	$28.25 \\ 46.75$	$32.50 \\ 47.25$
6 Cream 40 Whole Milk 4	$21.0 \\ 40.0$	$26.00 \\ 40.50$	$31.25 \\ 39.25$	$36.25 \\ 38.75$	41.25 38.50
7 Cream 30 Whole Milk 4	$\frac{29.0}{32.0}$	$36.00 \\ 30.50$	$\frac{43.25}{27.25}$	$50.00 \\ 25.00$	57.00 22.75
8 Cream 20 Whole Milk . 4	48.0 13.0	$58.50 \\ 8.00$	70.0 0.5		
Add to each above combination Unsweetened Condensed					
Skim Milk*	$24.00 \\ 15.0$	$18.5 \\ 15.0$	$14.5 \\ 15.0$	10.00 15.00	5.25 15.00
Gelatin	0.3	0.3	0.3	0.3	0.3
Total	100.0	100.0	100.0	100.0	100.0

^{*} Concentration, 3 to 1; contains 27% solids.

Note: The proportions given in columns a and b represent medium-fat ice creams commonly produced for soda fountain trade. These types of ice cream usually have a very smooth texture. The increased serum solids are derived chiefly from concentrated milk products. In some cases about 90% of the serum solids are added in the form of condensed skim milk, which means that approximately one-third of the mixture is condensed milk and one-fifth is cream testing 40 per cent fat. The cream flavor may be largely masked by the condensed-milk flavor, particularly if the latter has a pronounced cooked flavor. Consequently, the flavor will be improved by using either whole or skim milk with a minimum quantity of condensed skim milk.

The proportions given in columns e, d, and e represent ice creams with smaller additions of serum solids in the form of condensed skim milk, than those shown in columns a and b. It is believed that a small addition of serum solids to the higher fat products will improve the original texture, and in preventing deterioration of

texture during storage.

Table 3.—Amounts of Cream of Different Fat Content, Dry Skim Milk, and Fresh Skim Milk or Whole Milk Necessary for Making Different Types of Ice Cream

A. Solids		Type	s of Ice C	ream	
	a	\boldsymbol{b}	c	d	e
	Per Cent	Per Cent	Per Cent	Per Cent	Per Cent
Fat Serum Solids Sugar Gelatin Total Solids	10 11 15 0.3 36.3	12 10 15 0.3 37.3	14 9 15 0.3 38.3	16 8 15 0.3 39.3	18 7 15 0.3 40.3
B. Ingredients Per cent	C. Percentage		lient by W Cream Mix		Each Type of
	a	ъ	c	d	c
1 Cream 50 Skim Milk 0	20.00 60.00	$\frac{24.00}{37.00}$	$28.00 \\ 54.00$	$32.00 \\ 51.00$	$\frac{36.00}{48.00}$
2 Cream 40 Skim Milk 0	25 . 00 55 . 00	$30.00 \\ 51.00$	$35.00 \\ 47.00$	$40.00 \\ 43.00$	$\frac{45.00}{39.00}$
3 Cream 30 Skim Milk 0	$33.25 \\ 46.75$	$\frac{40.00}{41.00}$	$\frac{46.75}{35.25}$	53.25 29.75	$60.00 \\ 24.00$
4 Cream 20 Skim Milk 0	$50.00 \\ 30.00$	$60.00 \\ 21.00$	$70.00 \\ 12.00$	$80.00 \\ 3.00$	
5 Cream 50 Whole Milk 4	15.00 65.00	19.00 62.00	23.50 58.50	$27.57 \\ 55.43$	$32.00 \\ 52.00$
6 Cream 40 Whole Milk 4	$19.00 \\ 61.00$	$24.50 \\ 56.50$	$30.00 \\ 52.00$	35.25 47.75	$40.75 \\ 43.25$
8 Cream 20 Whole Milk 4	26.25 53.75	$33.75 \\ 47.25$	$41.50 \\ 40.50$	48.75 34.25	56.50 27.50
7 Cream 30 Whole Milk 4	$\frac{42.50}{37.50}$	$54.75 \\ 26.25$	$67.00 \\ 15.00$	$79.25 \\ 3.75$	
Add to each above com- bination					
Dry Skim Milk Sugar Gelatin	$\begin{array}{c} 5.00 \\ 15.00 \\ 0.3 \end{array}$	$\begin{array}{c} 4.00 \\ 15.00 \\ 0.3 \end{array}$	$\begin{array}{c} 3.00 \\ 15.00 \\ 0.3 \end{array}$	2.00 15.00 0.3	$1.00 \\ 15.00 \\ 0.3$
Total	100.00	100.00	100.00	100.00	100.00

Note: Dry skim milk is a very convenient form of serum solids to use in the manufacture of ice cream. Tests reported in U. S. Department of Agriculture Circular 179 have shown that the addition of dry skim milk will produce a medium grade ice cream equal to ice creams made with condensed milk. The principal criticisms of ice creams containing dry skim milk are usually due to the flavor imparted by this product. The formulas given in the above table will reduce this difficulty to a minimum by using as much whole and skim milk as possible in the preparation of the mixes.

No. 7			No. 8	
8% Fat—Cream, Skim Powder, 14%		Milk	8% Fat—Sweet Butter, Wh Skim Milk Powder, 14% S	
Sugar Gelatin Skim Milk Powder Cream (25%) Skim Milk	14 0.5 6 20 59.5	lb. lb.	Sugar Gelatin Skim Milk Powder Butter (84%) Milk (4%)	14 lb. 0.5 lb. 6 lb. 6.1 lb. 73.4 lb.

Table 4.—Amounts of Cream of Different Fat Content, Skim Milk, and Sweetened Condensed Whole Milk Necessary for Making Different Types of Ice Cream

A. Solids	Types of Ice Cream							
	a	ъ	c	d	e			
	Per Cent	Per Cent	Per Cent	Per Cent	Per Cent			
Fat Serum Solids	10 11	$^{12}_{10}$	$^{14}_{9}$	16 8	18 7			
Sugar	15	15	15	15	15			
Gelatin	0.3	0.3	0.3	0.3	0.3			
Total Solids	36.3	37.3	38.3	39.3	40.3			
B. Ingredients Per cent C. fat	Percentage	e of Ingred Ice (dient by W Cream Mix	'eight in E ture	ach Type of			
	\boldsymbol{a}	ъ	c	d	e			
1 Cream 50 Skim Milk 0	16.00 50.50	$20.68 \\ 49.72$	$25.6 \\ 49.2$	30.4 48.8	35.2 46.9			
2 Cream 40 Skim Milk 0	$20.00 \\ 46.50$	$26.00 \\ 44.40$	32.0 42.8	$38.0 \\ 41.2$	44.25 37.85			
3 Cream 30 Skim Milk 0	$26.66 \\ 39.84$	$34.7 \\ 35.70$	$\frac{42.7}{32.1}$	$50.7 \\ 28.5$	$58.7 \\ 23.4$			
4 Cream 20 Skim Milk 0	$\frac{40.00}{26.50}$	$52.0 \\ 18.4$	$64.0 \\ 10.8$	$\begin{array}{c} 76.0 \\ 3.2 \end{array}$	Marie Ma			
Add to each above com- bination	bination							
Sweetened Condensed Whole Milk*	25.00	20.00	15.00	10.00	5.00			
Sugar	4.5	6.6	8.7	10.8	12.9			
Water	4.0	3.0	1.5	10.0				
Gelatin	0.3	0.3	0.3	0.3	0.3			
Total	100.0	100.0	100.0	100.0	100.0			
* Contains 8% fat, 23% s	erum solid	s, and 42%	sugar.					

Note: Before using these formulas the manufacturer should be certain that the analysis of the sweetened condensed whole milk conforms to the analysis used in compiling this table.

	No. 9	
8% 1	Fat—Sweet Butter, Skir Milk Powder, 14% S	
Su	gar	14 lb.
Ge	latin	0.5 lb.
Sk	im Milk Powder	6 lb.
Bu	tter (84%)	9.6 lb.
Sk	im Milk	69.9 lb.
	No. 10	
8%	Fat—Sweet Butter, Milk Powder, 14%	
Su	gar	14 lb.
	latin	0.5 lb.
Bu	tter (84%	9.6 lb.
	im Milk Powder	12.6 lb.
Wa	ater	63.3 lb.
	No. 11	
	Fat—Cream and Milk im Milk 27% Solids, 1	
Su	gar	14 lb.
	latin	0.5 lb.

Condensed Milk	30	lb.
Cream (25%)	28	lb.
Milk (4%)	27.5	lb.

No. 12

8% Fat—Cream, Milk, Sweet Condensed Whole Milk, 14% Sugar.

Sugar	2.8 lb.
Gelatin	0.5 lb.
Sweet Condensed Milk	28 lb.
Cream (25%)	15 lb.
Milk (4%)	53.7 lb.

No. 13

8% Fat—Sweet Butter, Skim Milk and Sweet Condensed Skim Milk, 14% Sugar.

Sugar	2.8	lb.
Gelatin	0.5	lb.
Condensed Skim Milk	28	lb.
Sweet Butter	9.6	lb.
Skim Milk	59.1	lb.

Table 5.—Amounts of Cream with 50% of the Fat Added in the Form of Butter, Unsweetened Condensed Skim Milk, and Dry Skim Milk Necessary for Making Different Types of Ice Cream

A. Solids		Types of Ice Cream					
		a	Ъ	c	d	e	
		Per Cent	Per Cent	Per Cent	Per Cent	Per Cent	
Fat		10 11 15 0.3	12 10 15 0.3	14 9 15 0.3	$16 \\ 8 \\ 15 \\ 0.3$	$18 \\ 7 \\ 15 \\ 0.3$	
Total Solids		36.3	3 7. 3	38.3	39.3	40.3	
B. Ingredients Pe	er cent (C. Percentage	of Ingred	dient by W Cream Mix	eight in F	lach Type	of
		\boldsymbol{a}	ъ	\boldsymbol{c}	d	e	
1 Cream Unsweetened Condensed	40	12.50	15.00	17.50	20.00	22.50	
Skim Milk* Water		$28.00 \\ 28.40$	$\frac{34.00}{28.68}$	$30.00 \\ 28.96$	$26.00 \\ 29.25$	$22.00 \\ 29.53$	
2 Cream Dry Skim Milk† Water	40	12.50 10.80 55.60	15.00 9.67 53.01	$17.50 \\ 8.51 \\ 50.45$	$20.00 \\ 7.60 \\ 47.65$	$\begin{array}{c} 22.50 \\ 6.25 \\ 45.28 \end{array}$	
3 Cream Unsweetened Condensed	20	25.00	30.00	35.00	40.00	45.00	
Skim Milk* Water		$34.00 \\ 19.00$	$29.00 \\ 18.68$	$24.00 \\ 17.46$	$19.00 \\ 16.25$	$\frac{14.00}{15.03}$	
4 Cream Dry Skim Milk† Water	20	25.00 9.60 44.30	30.00 8.25 39.43	35.00 6.25 35.21	40.00 5.38 29.87	$45.00 \\ 6.00 \\ 23.03$	
Add to each above bination	com-	11.00	00.10	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	20.01	20.00	
Butter		6.10 15.00 0.3	$7.32 \\ 15.00 \\ 0.3$	8.54 15.00 0.3	9.75 15.00 0.3	$10.97 \\ 15.00 \\ 0.3$	
Total		100.0	100.0	100.0	100.0	100.0	
* Concentration	ratio, 3 t	to 1; contain	s 27% soli	.ds.			

Note: In the preparation of ice cream mixes with butter only the freshest and best grades of unsalted butter should be used.

No. 14		No. 16	
8% Fat—Cream, Milk, Evaporated 14% Sugar.	Milk,	10% Fat—Cream, Skim Powder, 12%	
Sugar 14	lb.	Sugar	12 lb.
Gelatin 0.5	lb.	Gelatin	0.5 lb.
Evaporated Milk (8%) 30	lb.	Skim Milk Powder	5 lb.
Cream (25%) 16	lb.	Cream (25%)	40 lb.
Milk (4%) 39.5	lb.	Skim Milk	42.5 lb.
No. 15		No. 17	
10% Fat—Cream, Whole Milk, Milk Powder, 12% Sugar.	Skim	10% Fat—Sweet Butte	r, Whole Milk,
Sugar 12	lb.	Skim Milk Powder, Sugar	12 % Sugar. 12 lb.
Gelatin 0.5	lb.	Gelatin	0.5 lb.
Skim Milk Powder 5	lb.	Skim Milk Powder	0 11.
		Skim Milk Fowder	6 lb.
Cream (30%) 26 Milk (4%) 56.5	lb.	Butter (84%) Milk (4%)	9 lb.

^{† 95%} solids.

Table 6.—Amounts of Cream of Different Fat Content, Unsweetened Condensed Skim Milk and Fresh Whole or Skim Milk Necessary for Making Different Types of Mixes Without Sugar

A. Solids		Types of Ice Cream				
		\boldsymbol{a}	ъ	c	d	e
		Per Cent	Per Cent	$\operatorname{Per}\operatorname{Cent}$	Per Cent	Per Cent
Fat		$10 \\ 11 \\ 15 \\ 0.3 \\ 36.3$	12 10 15 0.3 37.3	14 9 15 0.3 38.3	16 8 15 0.3 39.3	18 7 15 0.3 40.3
		C. Percentage	of Ingre		Weight in E	
		\boldsymbol{a}	\boldsymbol{b}	\boldsymbol{c}	d	e
1 Cream Skim Milk	50 0	$23.50 \\ 48.75$	$28.25 \\ 49.25$	$33.00 \\ 49.75$	$37.75 \\ 50.25$	42.50 50.50
2 Cream Skim Milk	40 0	$\frac{29.5}{42.75}$	$35.25 \\ 42.25$	$\frac{41.25}{41.50}$	$\frac{47.00}{41.00}$	$53.00 \\ 40.00$
3 Cream Skim Milk	30 0	$\frac{39.25}{33.00}$	$\frac{47.00}{30.50}$	$55.00 \\ 27.75$	$62.75 \\ 25.25$	$70.5 \\ 22.5$
4 Cream Skim Milk	$\begin{array}{c} 20 \\ 0 \end{array}$	58.75 13.50	70.50 7.00	$82.50 \\ 0.25$	***************************************	
5 Cream Whole Milk	$\frac{50}{4}$	$19.25 \\ 53.00$	$24.00 \\ 53.50$	$28.75 \\ 54.00$	$33.25 \\ 54.75$	38.00 55.00
6 Cream Whole Milk	40 4	$24.75 \\ 47.50$	$30.75 \\ 46.75$	$36.75 \\ 46.00$	42.50 45.50	$48.50 \\ 44.50$
7 Cream Whole Milk	$\frac{30}{4}$	$34.00 \\ 38.25$	$\frac{42.50}{35.00}$	50.75 32.00	59. 00 29. 00	67.25 25.75
8 Cream Whole Milk	$\frac{20}{4}$	55.50 16.75	69 . 00 8 . 50	$82.25 \\ 0.50$		
Add to each above bination Unsweetened Conde						
		$27.75 \\ 0.34$	$\begin{array}{c} 22.50 \\ 0.34 \end{array}$	$17.25 \\ 0.34$	$\begin{array}{c} 12.00 \\ 0.34 \end{array}$	$7.00 \\ 0.34$
Total		100.0	100.0	100.0	100.0	100.0

^{*}Concentration ratio, 3 to 1; contains 27% solids.

Note: Ice cream mixes made from the formulas in Tables 6 and 7 should not be confused with mixes containing sugar. For every 100 pounds of ice cream desired ase 85 pounds of mix and add 15 pounds of sugar. In case the manufacturer desires to use 1 or 2 pounds more or less of sugar, the basic formulas will not be materially changed.

No. 18			No. 19		
10% Fat-Sweet Butter, Sk	tim Milk	10% Fat-Sw			Iilk
Powder, Water, 12% S	ugar.	Powder, V	Vater, 12%	Sugar.	
Sugar	12 lb.	Sugar		12	lb.
Gelatin	0.5 lb.	Gelatin		0.5	lb.
Skim Milk Powder	6 lb.	Butter (84%)		12	lb.
Butter (84%)	12 lb.	Skim Milk P	owder	12	lb.
Skim Milk	69.5 lb.	Water		63.5	lb.

Table 7.—Amounts of Cream of Different Fat Content, Dry Skim Milk and Fresh Skim or Whole Milk Necessary for Making Different Types of Mixes Without Sugar

A. Solids	Types of Ice Cream				
	a Per Cent	b Per Cent	c Per Cent	$rac{d}{ ext{Per Cent}}$	ePer Cent
Fat	10 11 15 0.3 36.3	12 10 15 0.3 37.3	14 9 15 0.3 38.3	16 8 15 0.3 39.3	$ \begin{array}{c} 18 \\ 7 \\ 15 \\ 0.3 \\ 40.3 \end{array} $
B. Ingredients Per cent C.	Percentage		lient by W Cream Mix		lach Type of
	a	ъ	c	d	e
1 Cream 50 Skim Milk 0 2 Cream 40 Skim Milk 0	23.52 70.69 29.40	28.22 66.14 35.27	32.94 63.47 41.17	37.64 60.04 47.05	42.35 56.38 52.92
Skim Milk 0 3 Cream 30 Skim Milk 0 4 Cream 20	64.81 39.20 55.01 58.80	60.09 47.03 48.33 70.55	55.24 54.90 41.51 82.35	50.63 62.73 34.95 94.10	45.81 70.57 28.16
Skim Milk 0 5 Cream 50 Whole Milk 4	35.41 17.40 76.81	24.81 22.36 73.00	14.06 27.50 68.91	3.58 32.50 65.18	37.50 61.23
6 Cream 40 Whole Milk 4 7 Cream 30	22.27 71.94 30.77	28.60 66.76 39.60	35.00 61.41 48.50	41.50 56.18 57.50	49.00 49.73 66.25
Whole Milk 4 8 Cream 20 Whole Milk 4	63.44 50.0 44.21	55.76 64.26 33.10	47.91 78.75 17.66	40.18 93.25 4.43	32.48
Add to each above combination Dry Skim Milk*	$5.79 \\ 0.34$	4.64 0.34	3.59 0.34	$\frac{2.32}{0.34}$	$1.27 \\ 0.34$
Total* *Contains 95% solids.	100.0	100.0	100.0	100.0	100.0
No. 20		* .		No. 22	
10% Fat—Cream, Whole M Milk Powder, 14% Su			Fat—Swee		Whole Milk, & Sugar.
Sugar Gelatin Skim Milk Powder Cream (30%) Milk (4%) No. 21	14 lb. 0.5 lb. 4 lb. 26 lb. 55.5 lb.	Butt	tin 1 Milk Pow er (84%) 1 (4%)	der No. 23	14 lb. 0.5 lb. 4 lb. 9 lb. 72.5 lb.
10% Fat—Cream, Skim Milk, Powder, 14% Sugar	r .		Fat-Swe		Skim Milk, % Sugar.
Sugar Gelatin Skim Milk Powder Cream (25%) Skim Milk	14 lb. 0.5 lb. 4 lb. 40 lb. 41.5 lb.	Swee		vder (84%)	14 lb. 0.5 lb. 4 lb. 12 lb. 69.5 lb.

No	- 97
TA ()*	

10%	Fat—Sweet	Butter,	Skim	Milk	Pow-
, .	der. Wat				

Sugar	14 lb.
Gelatin	0.5 lb.
Butter (84%)	12 lb.
Skim Milk Powder	10.6 lb.
Water	62.9 lb.

No. 25

10% Fat—Cream, Milk Condensed Skim Milk (27%), 14% Sugar

(,0,,	
Sugar	14 lb.
Gelatin	0.5 lb.
Condensed Milk	18 lb.
Cream (30%)	28 lb.
Milk (4%)	39.5 lb.

No. 26

10% Fat—Cream and Milk, Sweet Condensed Whole Milk, 14% Sugar.

Sugar	6.8 lb.
Gelatin	0.5 lb.
Condensed Milk	18 lb.
Cream (25%)	27 lb.
Milk (4%)	47.7 lb.

No. 27

10% Fat—Sweet Butter, Skim Milk and Sweet Condensed Skim Milk, 14% Sugar.

Sugar	14	lb.
Gelatin	0.5	lb.
Butter (84%)	12	lb.
Condensed Skim Milk	16	lb.
Skim Milk	57.5	lb.

No. 28

10% Fat—Cream, Milk Evaporated Milk, 14% Sugar.

TEN Dugai.	
Sugar	14 lb.
Gelatin	0.5 lb.
Evaporated Milk	18 lb.
Cream (25%)	28 lb.
Milk (4%)	39.5 lb.

No. 29

12% Fat—Cream, Whole Milk, Skim Milk Powder, 12% Sugar.

· · · · · · · · · · · · · · · · · · ·	
12	lb.
0.5	lb.
2	lb.
41	lb.
44	lb.
	$12 \\ 0.5 \\ 2 \\ 41$

No. 30

12% Fat—Cream, Skim Milk, Skim Milk Powder, 12% Sugar.

Sugar	12	lb.
Gelatin	0.5	lb.
Skim Milk	3 _	lb.
Cream (30%)	40	lb.
Skim Milk	44.5	lb.

No. 31

12% Fat—Sweet Butter, Whole Milk, Skim Milk Powder, 12% Sugar.

	/0	
Sugar		12 lb.
Gelatin		0.5 lb.
Skim Milk Powder		3 lb.
Butter (84%)		10.8 lb.
Milk (4%)		73.7 lb.

No. 32

12% Fat—Sweet Butter, Skim Milk and Skim Milk Powder. 12% Sugar.

Skim wink rowder, 12%	ougar.
Sugar	12 lb.
Gelatin	0.5 lb.
Skim Milk Powder	3 lb.
Butter (84%)	14.3 lb.
Skim Milk	70.2 lb.

No. 33

12% Fat—Sweet Butter, Skim Milk Powder, Water, 12% Sugar.

Sugar	12 lb.
Gelatin	0.5 lb.
Butter (84%)	14.3 lb.
Skim Milk Powder	9.5 lb.
Water	63.7 lb.

No. 34

12% Fat—Cream, Whole Milk, Skim Milk Powder, 14% Sugar.

Sugar	14	lb.
Gelatin	0.5	lb.
Skim Milk Powder	2	lb.
Cream (25%)	41.5	lb.
Milk (4%)	42	1b.

No. 35

12% Fat—Cream, Skim Milk Powder, 14% Sugar.

Sugar	14	lb.
Gelatin	0.5	lb.
Skim Milk Powder	2	lb.
Cream (30%)	40	lb.
Skim Milk	43.5	lb.

No. 36

12% Fat—Sweet Butter, Whole Milk, Skim Milk Powder, 14% Sugar.

		•
Sugar	14	lb.
Gelatin	0.5	lb.
Skim Milk Powder	2.5	lb.
Butter (84%)	11	lb.
Milk (4%)	72	lb.

No. 37

12% Fat—Sweet Butter, Skim Milk, Skim Milk Powder, 14% Sugar.

1022-10	~~~~	•
Sugar	14	lb.
Gelatin	0.5	lb.
Skim Milk Powder	2.5	lb.
Butter (84%)	14.3	lb.
Skim Milk	68.7	lb.

FOOD PRODUCTS, BEVERAGES, FLAVORS 14				
No. 38		No. 45		
12% Fat—Sweet Butter, Skim Milk Powder, Water, 14% Sugar.		14% Fat—Sweet Butter, Whole Milk, Skim Milk Powder, 12% Sugar.		
Sugar Gelatin Butter (84%) Skim Milk Powder Water	14 lb. 0.5 lb. 14.3 lb. 9 lb. 62.2 lb.	Sugar Gelatin Skim Milk Powder Butter (84%) Milk (4%)	12 lb. 0.5 lb. 2 lb. 13.3 lb. 72.2 lb.	
No. 39	İ	No. 46		
12% Fat—Cream, Milk, Con Milk (27%), 14% S		14% Fat—Sweet Butter, Skim Milk, Skim Milk Powder, 12% Sugar.		
Sugar Gelatin Condensed Milk Cream (30%) Milk (4%)	14 lb. 0.5 lb. 16 lb. 35.5 lb. 34 lb.	Sugar Gelatin Skim Milk Powder Butter (84%) Skim Milk	12 lb. 0.5 lb. 2 lb. 16.7 lb. 68.8 lb.	
No. 40		No. 47		
12% Fat—Cream, Milk, Swe Whole Milk, 14% S	ugar.	14% Fat—Sweet Butter, Powder, Water, 12%	Skim Milk Sugar.	
Sugar Gelatin Sweet Condensed Milk Cream (25%) Milk (4%)	8.4 lb. 0.5 lb. 14 lb. 38 lb. 39.1 lb.	Sugar Gelatin Butter (84%) Skim Milk Powder Water	12 lb. 0.5 lb. 16.7 lb. 8.6 lb. 62.2 lb.	
No. 41	. 25:32 7	No. 48		
12% Fat—Sweet Butter, Sk Sweet Condensed Skir 14% Sugar.		14% Fat—Cream, Milk, Skim Milk Powder, 14% Sugar.		
Sugar Gelatin Sweet Skim Condensed M Butter (84%) Skim Milk No. 42	9.2 lb. 0.5 lb. ilk 12 lb. 14.3 lb. 64 lb.	Sugar Gelatin Skim Milk Powder Cream (30%) Milk (4%)	14 lb. 0.5 lb. 1 lb. 41 lb. 43.5 lb.	
12% Fat—Cream, Milk, Evaporated Milk, 14% Sugar. 14% Fat—Cream, Skim Milk Powder, 14% Sug				
Sugar Gelatin Evaporated Milk (8%) Cream (30%) Milk (4%)	14 lb. 0.5 lb. 20 lb. 30 lb. 35.5 lb.	Sugar Gelatin Skim Milk Powder Cream (25%) Skim Milk	14 lb. 0.5 lb. 1 lb. 56 lb. 28.5 lb.	
No. 43		No. 50		
14% Fat—Cream, Whole Milk Powder, 12%		14% Fat—Sweet Butter, Skim Milk Powder, 14	1% Sugar.	
Sugar Gelatin Skim Milk Powder Cream (30%) Milk (4%)	12 lb. 0.5 lb. 2 lb. 14 lb. 44.5 lb.	Sugar Gelatin Skim Milk Powder Butter (84%) Milk (4%)	14 lb. 0.5 lb. 1.1 lb. 13.3 lb. 71.1 lb.	
No. 44		No. 51		
14% Fat—Cream, Skim Mi Powder, 12% Sug Sugar Gelatin Skim Milk Powder	gar. 12 lb. 0.5 lb. 2 lb.	14% Fat—Sweet Butter, Skim Milk Powder, 14 Sugar Gelatin Skim Milk Powder	14 lb. 0.5 lb. 1.2 lb.	
Cream (25%) Skim Milk	56 lb. 29.5 lb.	Skim Milk	16.7 lb. 67.6 lb.	

No. 52

14% Fat—Sweet Butter, Skim Milk Powder, Water, 14% Sugar.

Sugar	14 lb.
Gelatin	0.5 lb.
Butter (84%)	16.7 lb.
Skim Milk Powder	7.6 lb.
Water	61.2 lb.

No. 53

14% Fat—Cream, Milk, Condensed Skim Milk (27%), 14% Sugar.

Sugar	14	lb.
Gelatin	0.5	lb.
Condensed Skim Milk	6	lb.
Cream (30%)	42	lb.
Milk (4%)	37.5	lb.

No. 54

14% Fat—Cream, Milk, Sweet Condensed Whole Milk, 14% Sugar.

Sugar	11.6	lb.
Gelatin	0.5	lb.
Sweet Condensed Milk (8%)	6	lb.
Cream (38%)	40	lb.
Milk (4%)	41.9	lb.
No. 55		

14% Fat—Sweet Butter, Skim Milk and Sweet Condensed Skim Milk, 14% Sugar.

, -		
Sugar	11.6	lb.
Gelatin	0.5	lb.
Condensed Skim Milk	6	lb.
Butter (84%)	16.7	lb.
Skim Milk	65.2	lb.
No. 56		

14% Fat—Cream, Milk, Evaporated Milk, 14% Sugar.

Sugar	14 lb.
Gelatin	0.5 lb.
Evaporated Milk	10 lb.
Cream (30%)	40 lb.
Milk (4%)	35.5 lb.

Fig Cream

For a 10-gal. finished ice cream.

45 lb. unflavored mix No. 10 can of solid packed pie figs ground fine in a food chopper is added while the mix is in the freezer.

Fig and Walnut Ice Cream

For a 10-gal. batch of finished product take 3 lb. canned pie figs, 2 lb. walnuts, run them through the fruit chopper, not too fine, and add the same as for strawberries. Use either English or black walnuts. The English are rather high in price.

The gelatin given in these formulas are .5 of a pound of high grade gelatin,

or you may use half good ice cream powder and half gelatin.

When mix is ready pasteurize the whole mix at 145 to 150° F., then viscolize or homogenize the whole mix while hot; cool to 40 or 50° F., age for 24 to 48 hours, then freeze.

Simple Ice Cream Mix

Cream (30%)	35.8 lb.
Milk (3.5%)	49.7 lb.
Sugar	14 lb.
Gelatin	0.5 lb.

100.0 lb. of mix containing 12.5% fat; and 33.4% total solids.

Complex Ice Cream Mix

Cream (30%)		41.7 lb.
Condensed Skim	Milk	15.3 lb.
Skim Milk		28.5 lb.
Sugar		14 lb.
Gelatin		0.5 lb.

100.0 lb. of mix containing 12.5% fat; and 37% total solids. Add 9¼ oz. standard vanilla extract to each 100 lb. of mix.

Preparing 20% Cream

To make 360 lb. of 20% cream use 160 lb. of 40% cream and 200 lb. of 4% milk.

Preparing 35% Cream

To make 360 lb. of 35% cream use 310 lb. of 40% cream and 50 lb. of 4% milk.

Chocolate Ice Cream

Milk		32	oz.
Sugar		16	oz.
Flour		2	oz.
Salt		1/8	oz.
Eggs		4	oz.
Cream		32	oz.
Vanillin		1/4	oz.
Unsweetened	Chocolate	4	07.

Heat milk and add flour, salt, and sugar. Stir thoroughly in double boiler for 20 minutes after batch is brought to a boil. After the mass thickens, add the beaten eggs and cook for 5 minutes longer with constant stirring. Cool, add cream which has been whipped into a stiff paste, and then add the flavoring. Add the melted chocolate, previously mixed with a litle sugar and warm milk to form a paste. Put in a refrigerator or pack in ice and salt until frozen.

Ice Cream Without Gelatin

Butter Fat	12	lb.
Sugar (Granulated)	12	lb.
Cerclose (Corn Sugar)	4	lb.
Milk Serum Solids	11.75	5 lb.
Vanilla Flavor	to	suit

Preventing Sandiness in Ice Cream U. S. Patent 1,940,109

By freezing and whipping air into icecream mix at such a rate that 30% of the water is frozen in less than 1 minute a smoother product than usual is obtained and one in which the milk solids may be increased with less likelihood of forming "sandy" ice cream.

Water Ices and Sherbets

The figures are given on the basis of 100 lb. of mix which is about 10½ gal.

Water Ice

Cane Sugar	25 lb.	
Corn Sugar	7 lb.	
Agar (3.2 oz. or 90.6 g.)	0.2 lb.	
Gum Tragacanth or Ga-		
lagum C (6.4 oz. or	0.4 lb.	
181.2 g.)	0.4 10.	
Water, Fruit, Fruit Acid, Flavor, and Color	67.4 lb.	

Overrun 20 to 25%. Total yield 13 gal.

Sherbet Using Milk

•		
Cane Sugar	25	lb.
Corn Sugar	7	lb.
Agar (3.2 oz. or 96.6 g.)	0.2	lb.
Gum Tragacanth or Ga-		
lagum C (3.2 oz. or		
90.0 g.)	0.2	lb.
Whole Milk	50	lb.
Water, Fruit, Fruit Acid,		
Flavor, and Color	17.6	lb.
Overrun 25 to 30%. Total	yield	13.5

Sherbet Using Cream Mix

gal.

Cane Sugar	25	lb.
Corn Sugar	7	
Agar (3.2 oz. or 90.6 g.)	0.2	lb.
Gum Tragacanth or Ga-		
lagum C (3.2 oz. or		
90.6 g.)	0.2	lb.
Ice Cream Mix, without		
Sugar or Gelatin	10-	lb.
Water, Fruit, Fruit Acid,		
Flavor, and Color	57.6	lb.

Overrun 25 to 30%. Total yield 13.5 gal.

Orange Water Ice (For 10 Gal. Batch)

Granulated Sugar	21	Ib.
Corn Sugar	7	lb.
Galagum C	3	oz.
Orange Juice (or Its Equiv-		
alent in Orange Flavor)	1	gal.

Citric acid to suit. Make up to 10 gal with water. Takes no overrun.

Orange Sherbet (10 Gal. Mix)

Cane Sugar	$22\frac{1}{2}$	lb.
Cerelose (Corn Sugar)	$7\frac{1}{2}$	lb.
Milk	4	gal.
Gelatin	11	OZ.
Orange Concentrate	4	oz.

Citric acid and color to suit. Make up to 10 gal. with water.

Cocoa Junket

Cocoa	2	OZ.
Boiling Water	4	oz.
Sugar	4.	oz.
Milk	32	OZ.
Junket Tablets	2	
Cold Water	1	oz.
Vanilla Extract	1/4	OZ.

Cook mixture of cocoa and water in double boiler for five minutes. Add sugar, stir until dissolved, and then add milk which has been previously preheated to 100° F. Add vanilla extract and heat to 120° F. Stir in junket tablets which are dissolved first in a little water. Pour into containers immediately, let stand until set.

Reworking Cream

For cream of poor quality mix equal parts of the cream and water and heat to 135° F. in a fore warmer. Condense in a vacuum pan until a volume equal to that of the original cream is obtained. Use 3 parts of cream to 1 part of water for cream that is of a slightly higher grade but that has off-flavors and odors. In this case fore warm and condense also until a volume equal to that of the original cream is obtained.

Composition of Mixes to Be Used in the Manufacture of Sweet Cream Cream Cheese

The most desirable cream cheese that has been manufactured by this method contains from 15 to 18% of dry skim

milk and 20% of butterfat in the final cheese mix.

The following mixes will make a very desirable cream cheese:

Formula No. 1

 Per hundred pounds:
 20
 lb.

 Butterfat
 20
 lb.

 Dry Skim Milk
 15
 lb.

 Gelatin (250 Bloom Test)
 0.4
 lb.

 Salt
 0.75
 lb.

Starter, 3 lb. (if cheese is for immediate consumption or 1 lb. if it is to be held in storage from 7 to 10 days prior to delivery to the consumer).

No. 2

 $\begin{array}{ccccc} \text{Per hundred pounds:} \\ \text{Butterfat} & 20 & \text{lb.} \\ \text{Dry Skim Milk} & 18 & \text{lb.} \\ \text{Gelatin (250 Bloom Test)} & 0.4 & \text{lb.} \\ \text{Salt} & 0.75 & \text{lb.} \\ \text{Starter} & 3 & \text{or } 1 & \text{lb.} \\ & & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & \\ & & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & \\ & & & \\ & & \\ & & & \\ & & \\ & & & \\ & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & \\ & & \\ & \\ & & \\ & & \\ & & \\ & \\ & &$

No. 3

 Per hundred pounds:
 25
 lb.

 Butterfat
 25
 lb.

 Dry Skim Milk
 15
 lb.

 Salt
 0.75
 lb.

 Gelatin (250 Bloom Test)
 0.4
 lb.

 Starter, 3 or 1 lb. as stated in No. 1.

It requires 7 to 10 days for a desirable mild acid flavor to develop in the cream cheese when only 1 lb. of starter is used in the cheese mixes. However, 3 lb. of starter is sufficient to develop the desired acidity by the end of the second day, providing a high quality starter is used in the cheese. If the cheese is to be held in storage for a period of approximately 30 days, 1 lb. of starter or a fraction thereof will develop the desired flavor. All equipment should be thoroughly sterilized prior to use and all ingredients must be of high quality.

The most desirable cream cheese is obtained when using No. 2, however either No. 1 or No. 3 furnishes a very desirable cream cheese.

The addition of dry skim milk, starter, salt and gelatin reduces the butterfat content of the resultant mix and sufficient fat must be added to the mix to replace the decrease in butterfat content by the addition of these ingredients. The addition of 1% of dry skim milk and other non-fat ingredients reduces the butterfat content of the finished cheese mix 0.26 of 1%. Therefore, in preparing a mixture that will furnish a butterfat content of 20% in the finished cheese when using 15% of dry skim milk, the cream from which the cheese is to be made must test 23.9% butterfat.

Cream Cheese (Geneva Method)
(Detailed Directions for 100-lb. Batches)

Acid Flavor

Add 5 lb. of dry skim milk to 93 lb. of sweet cream testing 40 to 42% milk fat. Then add 0.5 lb. of ground agar and 0.75 lb. of salt. The cream should be well agitated as the dry skim milk and agar are slowly added. Pasteurize at 180° to 185° F. for 5 minutes. Cool to 110° F. Add 0.75 lb. of commercial starter. Homogenize at 3500 lb. pressure using no strainer in the intake pipe line. The homogenizer should have been previously run with water at 160° F. or above. Place the cheese immediately into the final package. Chill in a refrigerator at 40° to a temperature of 70° F. and incubate for 12 to 24 hours to develop an acid flavor. Then chill to and hold at 40° F.

The acidity develops slowly and the rate of development is controlled by the percentage inoculation. Reducing the skim milk solids to 3% tends to soften the body of the cheese and increases the tendency towards some whey drainage and lower total acidity. The cheese may be softened by decreasing the homogenization pressure to 3000 lb. or firmed by increasing it to 4000 lb. More than 1 lb. of salt will retard and 1½ lb. almost check acid development. Cream color may be added before pasteurization, if desired, and it has the special advantage of reducing the intensification of color of cheese exposed to the air.

Consideration has also been given to the omission of starter and the securing of the desired acid flavor from Neufchatel, cottage, or Neufchatel cream The process itself presented no cheese. special difficulties (even cottage cheese could be homogenized in the cold or warm cream at 100 lb. pressure) and the mixture was treated in the regular way. About 50% of these acid cheeses is required to impart a very mild acid flavor to the finished product; or a product such as that made from an enriched milk by the cottage cheese process could be homogenized alone. The process is somewhat complicated and the flavor of the finished cheese is very mild, but it has

excellent keeping quality.

The homogenizer may be a source of microbial contamination and may chill the first material passing through it. For these reasons the hot water rinse just before use is always essential. The cream mixture was strained through a coarse strainer with approximately ½6 inch openings and the strainer to the homog-

enizer was always removed from the pipe line to permit an even flow of the cream mixture. Short pipe lines are very desirable to reduce mechanical losses.

The hot cheese may be transferred with a filling machine or by hand to 3-or 5-lb. lined boxes for bulk sale. The usual mayonnaise jar filling machine can be used for filling jars, but some difficulty may be encountered in making the small tin foil or cellophane-wrapped 1- to 4-oz. packages. These packages are made from the cold cheese by molding into proper size with a machine or by cutting into the proper size with a remodeled butter cutter. Some ingenuity must be used in the details of placing the cheese in the package.

Ripened Cheese Flavor (Cheddar and Roquefort)

Add 5 lb. of dry skim milk to 69.25 lb. of sweet cream testing 40 to 42% of fat. Then add 0.75 lb. of common salt. (The agar is not essential in this cheese, but it improves slicing qualities.) The cream should be well agitated as the dry skim milk is slowly added. Remove paraffin, cheesecloth, or other coating from the surface of 25 lb. of well-ripened American cheddar cheese and grind or slice the cheese. Cheese color appears to be desirable for cream cheese of the cheddar flavor to give the cream the usual cheddar cheese color.

For Roquefort flavor use 79.25 lb. of sweet cream, 5 lb. of dry skim milk, 15 lb. of Roquefort cheese and 0.75 lb. of common salt. The entire mixture should be pasteurized at 160° or at 180° F. for 5 minutes, depending upon the keeping quality desired. Homogenize at 3500 lb. pressure, the machine having been previously run with hot water. Place the hot cheese directly into the final package and immediately store at 35° to 40° F. in the refrigerator.

Less Roquefort cheese is generally required as a flavor than is the case for American cheddar. Many persons who object to the flavor of Roquefort cheese consume large helpings of Roquefort cream cheese. Other varieties of cheese may be used, but investigations have been limited to the two varieties mentioned.

The ripened cheeses readily soften and disperse in the cream when the temperature exceeds 145° F. No necessity of using an emulsifying salt was ever encountered, but tests demonstrated that these salts, such as di-sodium phosphate and sodium citrate, could be used in

limited amounts without interfering with the process.

Other Food Flavors

Coarsely ground sweet pickle relish (onion flavor is undesirable), pimiento, olive and nut, pineapple, and other food flavors may be used. Add 5 lb. of dry skim milk, 0.5 lb. of ground agar, and 0.75 lb. of salt to 73.5 lb. of cream testing 40 to 42% of fat. The cream should be well agitated as the dry skim milk and agar are added. Pasteurize at 180° to 185° F. for 5 minutes. Homogenize at 3500 lb. pressure, the machine having been previously run with hot water. Stir the flavoring material, 20 lb. is about right for most foods, directly into the hot cheese. Place in the final package and store immediately in the refrigerator at 35° to 40° F.

In some instances there may be an excessive quantity of juice. This can be mixed in the cream just before homogenization, but if the acidity of the juice is high the cream mixture may be previously cooled to 120° to 140° F. before adding the juice and the homogenization pressure reduced to prevent excessive fat clumping and coagulation. If the body is somewhat soft the dry skim milk may be increased to 7 lb.

Most fruit flavors did not blend well with cream cheese, but tart spicy flavors are generally satisfactory.

O. and N. Cream Cheese (Marquardt)

Standardize milk to 10% of fat, then pasteurize at 160° F. for 30 minutes; and homogenize at 2500 lb. pressure and at 120° F.

Cool the batch to 72° F., and add 0.2% of commercial starter and 15 cc. of remet per 1000 lb. of milk. On the following day drain and salt as in the making of old style cream cheese and analyze for fat.

Mix the cheese prepared in the above manner with 40% cream to obtain the desired cheese fat content. This may be 27, 30, 35 or 40%. Then add 0.1% of gum and 5% of 40% sour cream. Add enough salt to have 0.75% in the finished cheese. Heat this entire mixture to 160° F. and homogenize at 120° F. and 3000 lb. pressure.

Bel Paese Cheese (Farrar)

Use raw milk containing 3 to 4% of fat. Add ½% of lactic culture, and an equal amount of S. thermophilus culture when available. Set the milk at 107° F. with rennet at the rate of 8 oz. per 1000

lb. of milk. The curd is cut after 15 minutes. Then part of the whey is drawn, and the cheese curd is dipped

rapidly into the molds.

The cheese should drain on reed mats for 6 hours, being turned frequently. It is desirable to have the room at 80° F. The cheese can be made in brick molds or circular ones 8 inches in diameter. The cheese should be of a thickness when finished so that it will weigh 3 to 5 lb.

The cheeses are salted by submerging in 20% salt brine at 50 to 60° F. for 18

to 24 hours.

The cheeses after drying are placed in a curing room at 40° F, with a relative humidity ranging from 85 to 90° F.

After curing the cheeses are wrapped and packed so as to avoid evaporation. This is exceedingly important. The cheeses cure in 6 to 12 weeks, depending upon the quality of the milk used.

Semi-Soft Cheese (Marquardt)

Use raw or pasteurized milk testing 3.5% in fat. Use 1 oz. cheese color per 1000 lb. of milk. Then add 44% of commercial lactic culture and 44% of S. helveticus culture and heat to 87° F. In about 2 hours the acid will increase .02 to .04 in the milk. Then dilute 8 oz. of rennet in cold water and add at this rate for each 1000 lb. of milk.

The milk should set for 30 minutes, and, 30 minutes after cutting it is dipped rapidly into brick or round molds. It is pressed with 10 lb. pressure for 8

hours.

After 24 hours the cheese is rubbed lightly with salt, and then placed in a brine for 24 to 48 hours. The brine is made by dissolving 18 lb. of salt in 82 lb. of water.

The cheese is cured at 53-57° F. for a short time, about 3 weeks. It is then

placed in storage at 40° F.

Each cheese should weigh from 3 to 7 lb.

Walter Price Rapid Cottage Cheese Method

Pasteurize skim milk. Cool to 90° F. and add 5% of culture. Acid development of 0.5% will require only 5 hours. Finish making cheese according to standard procedure.

Note: Setting at 72 to 85° F. requires 12 to 18 hours for 0.5% of acid to

evelop.

Propagating lactic culture:

Select good grade of skim milk. Pasteurize to 180° F. for 1 hour. Cool to 72° F. Add 1% of culture from another

culture. Incubate at 72° F. for 12 hours. Place in 40° F. room until ready for use.

Selecting natural culture:

Place 6 qts. of raw skim milk into a 72° F. incubator. After 12 hours select those having a firm curd. Select of the firm curd samples the one having best flavor. Use this as a propagating culture for future batches. Always inoculate from a day old culture.

Developing a commercial culture (Strep. Lact.):

Pasteurize skim milk to 180° F. for 1 hour in quart bottles. Cool to 72° F. and add a few drops of culture from a commercial culture. Incubate for 12 hours. Repeat pasteurization of a fresh batch of skim milk; and inoculate 1% from above culture. Repeat for 3 days, always using the culture just previously developed. After this period the culture is ready for use in cheese, butter, or cultured milk manufacture. Cultures should be transferred daily, and used for 3 weeks or a shorter period.

Developing Special Cultures (Bac. Bulgaricus of Lacto bacillus Acidophilus).

Follow above procedure for commercial cultures.

Incubate at 98° F.

Goats' Milk Cheese

Heat fresh milk to 88° F. Add 25 drops of rennet for each 10 lb. of milk. Before adding rennet dilute it in 20 times its volume of water. Cut in cubes 1 in. square after 45 minutes. Allow to stand for 5 minutes, then dip into molds after stirring gently for 5 additional minutes.

The forms are made of 3X tin; they are 4½ in. in diameter, and 5 in. high. Each form has 5 rows of holes, the holes being 1 in. apart and ½ in. in diameter.

The cheese curd is not disturbed until it is sufficiently matted. It is then turned frequently. It remains in the hoops for 30 hours at 70° F. It is then rubbed with salt and placed in a curing room at 60° F. with a high humidity. The cheese should be wiped freely and turned. After 6 weeks they are ready to package. Each cheese weighs ½ lb. and requires 4½ lb. of milk. The cheese is white and has an agreeable flavor at 6 to 10 weeks.

Hokah Sage Cheese

To 69¼ lb. of 40% fat content cream add 5 lb. of dry skim milk. Then add ¾ lb. of common salt and a like amount

of agar agar (ground or powdered). Slice and grind 25 lb. of well cured cheddar cheese into the mixture and stir while heating the batch to 160 to 180° F. Hold at this temperature for 2 minutes and cool to 140° F. Then add 1 to 3 cc. of oil of Sage, Dalmatian. It should be diluted in a pint of water and then mixed into a gallon of the cheese mixture which in turn is mixed into the entire batch. The mixture is then homogenized at 3500 lb. pressure, the machine having been previously run with hot water. If the minimum amount of sage oil is used 1/2 oz. of sage leaves, Salvia officinalis, may be added to the batch after homog-In using the leaves great care must be exercised in pulverizing them and removing stems and coarse leaves. Thorough incorporation is an essential. Extensive trials have indicated the desirability of using the oil of sage

The cheese should be packaged while hot, and stored at 35 to 40° F.

Cheese Pikante (Marquardt	Meth	od)
Roquefort Cheese	20	Ib.
Cheddar Cheese	20	lb.
Camembert Cheese	20	lb.
Salt	1/4	1b

Add small quantities of black pepper, cayenne pepper, paprika, and grind through a fine grinder. The addition of 2 to 4% of Sauterne Wine improves the Pikante. Grind with products at 70° F., package and store at 32 to 40° F.

New York Style Sage Cheese

The regular method for making cheddar cheese is followed. At the start 100 lb. of milk for colored curd is used for each 1000 lb. of milk. The small batch of milk is colored green. Both batches are made alike. At cheddaring time the curds of both batches are mixed and matted. Before pressing oil of sage, Dalmatian is atomized over the curd at the rate of ½ oz. per 1000 lb. of milk used.

The green color is prepared by soaking green corn, green oats, or alfalfa in water, grinding, and pressing in a cider press. The color must be prepared fresh each day. The amount to add to the small batch of milk depends upon the intensity of color desired.

Some manufacturers prefer to add the oil of sage to the milk before making

The above method appears to be the one most commonly used. Other methods

have been described but produce less satisfactory results.

Ricotta Cheese (Marquardt)

Heat whey to 190° F. as it is drawn from the cheese vat. Then add sour whey until albumin flakes are like snowflakes. Stop heating when albumin collects on top of whey. Drain in molds or bags. The cheese after draining is surface salted and ready for use.

The sour whey used should have 1% of acid. It may require a Bulgaricus culture to achieve this. To flake out the albumin about 10% of sour whey must be added to the sweet whey. When whey only is used and drained in bags the cheese is called mejette.

Commonly 10% of skim milk is added to the sweet whey to increase the yields.

Hoops used as molds should be 5 inches in diameter and 9 inches high and perforated. If the molds are completely filled with moist cheese with a strainer dipper the cheese resulting will be 7 inches high. The cheese is rubbed with salt and returned to the hoops for 2 hours after the draining period over night without pressure. The cheese should be dried in a room at 110° F. and wrapped in paper and placed in storage.

Maroni Cheese (Marquardt)

This is made by using the Ricotta method substituting whole milk for skim milk and adding 10%. It is molded in hoops 8 inches in diameter and 10 inches high, giving a finished cheese 7 inches high. Ricotta Gras is also the name for the whole milk-whey combination.

Sapsago Cheese

This cheese is made principally in Glarus, Switzerland, from sour, skim milk of cows. It is known also as Schabzieger, Glarnerkase, and Krauterkase. It is claimed to have been made in the thirteenth century; the authentic history at least dates back to the fifteenth century. Sapsago is a small, hard, green cheese flavored with the leaves of a species of clover; it is shaped like a truncated cone, 4 inches high, 3 inches in diameter at the base, and 2 inches at the top. This cheese is imported to some extent into the United States under the name of Sap Sago.

The skim milk from which this cheese is made is not allowed to become sour enough to coagulate on heating, as it would make too hard a curd. The milk, when it has reached the right acidity, is

heated to the boiling temperature while being stirred. Cold buttermilk is then added, as is also some whey having a high percentage of acidity. The material coagulating on the surface is skimmed off. The milk is then stirred, while sufficient acid whey is added to precipitate the casein. When too little whey is used the curd is too soft, and when too much is used it is too hard. The curd is dipped with a skimmer and spread out to cool and then put into boxes and allowed to drain and ferment. The box is kept at a temperature of above 60° F., and pressure is applied by weighting with stones. Ripening is allowed to continue from three to six weeks. If the temperature of the room is too high or if sufficient pressure is not applied, too rapid and strong fermentation results. The curd is used for making the finished product, but the cheese is seldom finished where the curd is made. The curd is ground in a mill, and for every 100 lb. of cheese there is added 5 lb. of salt and 25 lb. of dried Melilotus caerulea, an aromatic clover which is grown in the Canton of Schweiz for the purpose. The ground material is worked up into a dough and is forced into molds lined with linen cloth and the name of the manufacturer is stamped on the large end. The mold is then emptied and refilled. The cheeses are dumped promiscuously into a large cask holding about 200 lb. A comparatively small quantity is shipped into this country. It sells at a low price and is usually grated.

Red Cheese Rind Color

Formula No. 1

Sudan 4 dye is dissolved in equal parts of 70% alcohol and acetone, or

No. 2

Tournesol, Fuchsin, or Bordeux Red dissolved in water (distilled water is preferred), or

No. 3

Iron Oxide, known also as Berlin Red or English Red made into a paste with a heavy oil.

The intensity of the color can be varied by changing the amount of the coloring substance.

Apply to outside of cheese.

Cheese, Ice-Cream and Salad Stabilizer U. S. Patent 2,007,218

Locust Bean Gum	65	oz.
Irish Moss, Powdered	35	OZ.
Karaya Gum	15	oz.

When used in the preparation of cream cheese, the undiluted mixture of the three ingredients mentioned above is added at the time that the curds are mixed with the cream in the usual procedure for the manufacture of cream cheese, and in the proportion of about one-half of 1% by weight on a wet basis. The material is heated to about 165° F., homogenized, and then packed hot.

In ice cream it is used diluted with sugar, in the preferred proportion of one-half of 1% on a wet basis, the stabilizer acts to prevent crystallization of ice particles and thus insures a fine, smooth texture and a body which will hold up under severe shocks, such as are encountered in transportation and handling. The use of it in ice cream also usually results in more rapid freezing, especially in old-style freezers.

Cheese Emulsifiers

U. S. Patent 1,940,031

1-4% of either of following are used: Sodium Mucate Sodium Lactate

Preservation of Rindless Cheese British Patent 434,374

Bacterial action on surface of rindless cheese is prevented by treatment with following prior to heating to 65° C.

Hydrogen Peroxide (35%) 0.3%

Low fat content cheese is heated to 65° C. The peroxide is added, mixed and later heated to 80° C.

Brandy Cheese

Use regular cheddar cheese, preferably an entire small cheese with the surfaces scraped clean, and allow to dry at room temperature for 2 to 4 weeks. Then place cheese in clear water at 40 to 80° F. for several days.

The cheeses are then placed in a mixture of brandy and high grade vinegar for several days. The brandy may be mixed in equal parts or less with the vinegar. Three per cent of salt should be added with a liberal addition of pepper to the brandy-vinegar solution.

Sour Cream

To 20% cream add 2 to 3% skim milk powder. Heat slowly to 120° F. to dissolve the powder and follow this by pasteurizing at 145° F. for 30 minutes. Cool to 70 to 72° F., and add 3 to 5% of

good starter, thoroughly broken up. Dilute 20 drops of commercial rennet extract in about ¼ glass of water and add this to 100 lb. of cream, agitating it thoroughly to distribute the rennet. The rennet helps to form a thick curd and the cream may curdle in a relatively short period. However, you should hold it over night at the ripening temperature of 70 to 72° F. to develop the desired acid flavor. Follow this by breaking up the curd while cooling to 40° F. and hold at this low storage temperature.

Infants' Milk, Synthetic

Sugar	40	g.
Soya Bean Powder	125	g.
Lactose	30	
Peanut Oil	20	
Dextrin	20	g.
Egg Yolk, Liquid	50	ğ.
Calcium Lactate	6	
Salt	2	g.
Stir in water before use.		

Soya Bean Vegetable Milk

If the dried beans, preferably yellowseeded varieties, are soaked for a few hours, then finely crushed and boiled for about 30 minutes in the proportion of 3 parts of water to 1 part of mash, a milky emulsion is obtained which is very similar in appearance and properties to animal milk. This liquid, separated out by means of a very fine sieve or cloth strainer, is the Soya Bean or vegetable milk used so extensively in China. Soya bean meal after the oil is extracted or whole soya bean meal may be utilized quite as well as the whole bean. In the absence of animal milk, soya bean milk is used extensively in the fresh state and as the basis of various kinds of vegetable cheeses in oriental countries. Soya bean milk in the form of a powder is a commercial product in some European countries, and in parts of the United States it has been used in special feeding cases. The milk can be used successfully in numerous preparations, such as breads and cakes, in creaming vegetables, in milk chocolate, and in custards.

After separating the liquid from the solid material, the residue is still very rich in nutritive substances and can be dried and used for cattle feed or made into flour for human food.

Soya Bean Curd

The addition of magnesium or calcium salts or of rennet or lactic acid to soya bean milk when hot precipitates some of the protein, forming a grayish white curd which settles out, leaving a yellowish water liquid. This curd, after being drained and pressed, represents bean curd of tofu, which is extensively eaten and forms the basis of numerous fermented, smoked, and dried cheeses in China and Japan. Bean curd is made fresh daily and is a staple article of diet among oriental peoples. In many cities of the United States having a large oriental population fresh bean curd may be found in the Chinese and Japanese markets.

Dry Mix for Making Chocolate Milk in Dairies

Cocoa	1.75	lb.
Cane Sugar	7	lb.
Agar, Powdered	0.14	lb.
Vanillin	0.003	lb.
Salt	0.025	lb.

Mix the above ingredients well and add to each gallon of milk in the pasteurizer at 185° F. Agitate and hold for ½ hour.

Cocoa Malt Powder

Cocoa Powder	23 lb.
Fine Granulated Sugar	70 lb.
Malt Powder, Mild Flavor	20 lb.
Skim Milk (Soluble)	14 lb.
Sodium Bicarbonate	2 oz.
Salt	8 oz.
Vanillin	1/2 OZ.
Vanilla Extract	1/2 oz.

Mix ingredients thoroughly and pass through a coarse sieve. This mixture can be packaged in cans, glass containers, or in 1¼ oz. envelopes for individual use.

Stable Chocolate Milk U. S. Patent 1,989,758

In carrying out the process of making the milk starch emulsion, the chocolate, sugars (when the latter are used), starch, and the gum may be introduced, as dry substances, into the milk, thoroughly mixed, and the mixture heated to a temperature of 170° to 200° F., or higher if desired-although this is not necessary—in place of temperatures approximating 240° F. heretofore recommended, for periods from 20 to 30 minutes, more or less. Preferably, however, a syrup is first made of the chocolate and sugar, and this syrup, together with a preformed mixture, in proper proportions, of the starch and gum, added to the milk and the final mixture agitated and heated as described.

As a matter of convenience to the beverage manufacturer, and in order to insure correct proportions between starch and gum, the starch and gum may be compounded together and the compound delivered to the beverage manufacturer.

In making the compound the agaragar, for example, is prederably ground dry and screened to the same degree of fineness as the starch and is then thoroughly mixed with the starch in the proportions indicated by the specific examples given below. In such a mixture the agar-agar, although very small in quantity, approximately from 1 to 20 parts of agar to 100 parts of starch, will remain evenly distributed in the starch. It will not sift out. This novel mixture will disperse in the chocolate vehicle much more easily than if the ingredients were introduced into the liquid as separate substances. If the agar is not finely ground it will swell instead of dissolving, particularly at the low temperatures preferably used in compounding, with consequent loss of stabilizing

The following examples of typical mixtures, with preferred percentages of the ingredients, will serve to illustrate the character of the present invention. The percentages are by weight.

Formula No. 1

roimua no. r	
Milk	90.48
Cane Sugar	4.82
Dextrose (Cerelose)	2.41
Cocoa (High Grade, Dark) Raw Tapioca Starch (Scott	1.27
Raw Tapioca Starch (Scott	
Test 150)	1
Agar-Agar	0.02

Any suitable sugars may be used in the suspension or in the dry product or the sugar ingredient may be omitted if desired. The amount of the sugar in-gredient may be varied to any extent. For any usable quantity the sugar does not add to the viscosity of the beverage. The amount of cocoa or chocolate may also be varied. The matter of taste or of economy will govern any increase or decrease. As much as 2.5% of cocoa may be used without changing the percentage of starch or gum. The starch ingredient may be increased to 2 or 3%. Experience goes to show that 1% is near the critical lower limit. More than 2 or 3% gives too high a viscosity and is likely to give a distinct starch taste to the product. The agar-agar may be varied in amount from about 0.01% to 0.2%, but at the upper limit there is a strong tendency to segregation in jellylike lumps.

No. 2

90.78
4.06
2.03
1.673
1.433
0.024

The first four items may be varied as indicated in No. 1.

The same quantity of modified corn starch may be used in place of the specified raw corn starch. The amount of corn starch may vary between 1 and 2%. Where raw corn starch is used the lower limit of the gum quantity should not be quite as low as in No. 1.

No. 3

Milk	91
Cane Sugar	4.07
Cerelose (Corn Sugar)	2.03
Cocoa	1.676
Wheat Starch (Scott Test 85)	1.2
Gum	0.024

The variations may be substantially the same as with No. 1.

The time of cooking with the raw corn starch should be ordinarily 25 to 30 minutes; with the modified corn starch 20 to 25 minutes; with the tapioca and wheat starches about 20 minutes.

Chocolate-Flavored Milk

In this improved formula use:

THE CHILD THE PLOYER TOTAL	abo.	
Cocoa	20	lb.
Sugar	90	lb.
Skim Milk	90	lb.

To the above syrup add 2000 lb. of milk; heat to 143½° F. and hold for 30 minutes. Homogenize the mixture at 2000 to 3000 lb. pressure while hot.

Cool and bottle.

Non-Settling Cocoa Milk

Tion comme	00000	212,1712	
Cocoa Powder		6	oz.
Sugar		28	oz.
Sodium Alginate		1.	oz.
Milk		15	qt.

Mix together the cocoa, alginate, and sugar. Heat the milk to 160° F., add the dry mixture slowly with constant stirring, for thirty minutes. Cool the batch to 45° F. and hold for two hours before bottling in sterilized bottles. The cocoa powder can be of any fat percentage from 10 to 25%. The milk can be either whole milk or skim milk, or any

mixture of each. Additional flavoring ingredients such as vanilla, malt, caramel, etc., may be added.

Boiled Cocoa Frosting

Sugar	16	oz.
Salt	1/16	oz
Water	16	oz
Vanilla Extract	1/4	oz.
Dairy Butter	3/4	OZ.
Cocoa	3	oz.
Corn Starch	1	OZ.

Mix sugar, cocoa, and salt together, then add slowly 8 oz. of boiling water and when all the water has been added bring mix to a boil. Make a pre-mix of corn starch and cold water, then add to the above mix and again bring to a boil. Continue boiling with low flame, until the frosting has become thickened which usually requires 3 or 4 minutes. Remove from flame, add butter and vanilla extract, beat well, allow to set and cool.

Chocolate Filling

Milk	-	8	oz.
Sugar		2	oz.
Flour		11/2	oz.
Salt		1/16	oz.
Whole Eggs		2	OZ.
Vanilla Extract		1/4	oz.
Unsweetened Chocolate		1	oz.

Heat milk to boiling point, add sugar, flour, and salt, stirring thoroughly. Cook for fifteen minutes, add eggs slightly beaten, cook for 5 minutes longer. Add flavoring and unsweetened chocolate, and 1 oz. powdered sugar and stir.

Chocolate Mocha Frosting

Powdered Sugar	1½ lb.
Hot Coffee	3 oz.
Unsweetened Chocolate	2 oz.
Butter	¾ oz.

Moisten the sugar with coffee, blend the chocolate with dairy butter. Mix the two blends together and beat until smooth.

Chocolate Icing

Unsweetened Chocolate	e 2	OZ.
Water	4	OZ.
Sugar	16	OZ.
White of Eggs	2	OZ.
Vanilla Extract	1/16	OZ.

Warm water, then add powdered sugar, cook to approximately 216° F. until the mix threads well on the end of a spoon. Stir in the well-beaten white of eggs, then add melted chocolate and vanilla

and stir thoroughly to proper consistency.

Bailed Marshmallow for Topping Formula No. 1

No. 1

Granulated Sugar	3 lb.
Glucose	12 oz.
Water	1 pt.
Boil to 240° F.	
No. 2	

Egg Whites Granulated Sugar	1¼ pt. 8 oz.	
No. 3		
Water	4 oz.	

Vanilla Extract ½ oz.

XXXX Sugar 4 oz.

Method: Set contents of No. 1 into copper kettle, dissolve well together, and place over moderate fire. Set contents of No. 2 in small 12-quart machine kettle. Warm contents of No. 3 in small bowl and thoroughly dissolve.

When contents of No. 1 reach 225° F., start machine going with No. 2 on high speed. Also see to it that sides of copper kettle are kept clear of sugar crystals, by washing sides of kettle with water and brush.

The meringue content of No. 2 should be ready about the same time that the boiling content of No. 1 reaches the degree of 240° F.

With the meringue ready, and the boiled sugar at 240° F., pour the boiled sugar on to meringue slowly in thin stream (this is important). Let the machine run on high speed during this operation.

Now add dissolved contents of No. 3 to the mass, and continue whipping on high until a fine bodied smooth meringue is obtained.

Formula No. 2 (Quicker Method)

No. 1

Egg Whites	1 pt.
Granulated Sugar	1 Îb.
XXXX Sugar	8 oz.
Tapioca Flour	½ oz.
No. 2	
Glucose	4 oz.
Water	4 oz.
Gelatin	16 02

Method: Dissolve contents of No. 1 all together over double boiler and heat to 120° F. Keep contents stirred with wire hand whip. Now set kettle in machine

Vanilla Extract

and, with wire whip attached, beat on high speed. Immediately dissolve contents of No. 2 by warming, until all are dissolved together, then pour into machine and continue whipping until a fine meringue is obtained.

Whipped Cream for Baker's Topping

The cost of whipping cream and the fact that it will not stand up alone for very long makes its use almost prohibitive.

Fortified Whipped Cream

Cold Water	5	qt.
Meringue Powder	6	oz.
Sugar	4	lb.
Salt	1	oz.
Starch	14	oz.
Gelatin	3/4	oz.
Vanilla Extract	1	oz.
Heavy Cream	1	qt.

In the machine put 1 qt. of water, the meringue powder and 3 lb. of sugar and whip to just peak (not stiff). Put 3 qt. of water, the remaining sugar and the salt into a kettle and bring to a boil. Dissolve the starch and gelatin in the remaining water, add to the boiling mass and stir until it is thick and clear. Blend the two mixtures carefully with a wire whip and put in the refrigerator until When ready to use, put the mixture into a clean bowl and smooth down with a wire beater. Do not beat. Bring the whipping cream up to about three-fourths stiff, pour it over the boiled mixture, and fold together only until the cream is well incorporated and the mass is smooth.

This should make topping enough for 30 to 40 9-inch pies.

Baker's Pectin Glaze

Pectin	4 oz.
Sugar	8½ lb.
Water or Fruit Juice	21/2 qt.
Phosphoric Acid	$2\frac{1}{2}$ oz.

Mix the pectin with some of the sugar. Bring the liquid to a boil and add to the sugar-pectin mixture. Take off the fire and stir until the sugar is thoroughly dissolved. When this has been done add the remaining sugar, stirring in the meantime. Allow the liquid to cool, then add acid.

Coat the berries as much as possible and they will not have a chance to "bleed" and thus soak through into the cake itself. If desired the berries may be dipped into the glaze before they are applied to the cake and the remaining pectin poured over them so they are nicely coated.

Baking Powder

Sodium Acid Pyrophosphate	42	oz.
Sodium Acid Carbonate	30	oz.
Maize (or Rice) Starch	28	oz.

Stable Baking Powders German Patent 599,493

Formula No. 1

Cream of Tartar Tartaric Acid Sodium Bicarbonate Wheat Flour Carbamide Magnesium Peroxide	44 g. 6 g. 27 g. 20 g. 1.5 g. 1.5 g.
No. 2	
Calcium Biphosphate Sodium Bicarbonate	34 g.

Wheat Starch Powder

Carbamide

1.5 g. Magnesium Peroxide 1.5 g.

40

1,0, 9		
Sodium Acid Pyrophosphate	44	g.
Sodium Bicarbonate	32	g.
Maize Starch Powder	22	
Carbamide	1	g.
Magnesium Peroxide	1	g.

15 g. of above baking powders are used for 500 g. flour.

Soya Bean Flour Bread

Formula No 1

1.0111	idia 140.	.1.			
Soya Flour			6	5	1b.
Wheat Flour			26	0	lb.
Sugar			1	0	lb.
Salt				5	lb.
Yeast			1	5	lb.
Shortening			1	5	lb.
Water (Variabl	e)		21	0	lb.
Mix 3 minutes,	ferment	at	90°	F	۲.
First punch			45	n	in.
To bench			15	n	in.
\mathbf{Proof}			45	n	nin.
Bake			30	n	nin.
Temperature of	Oven	4	45°	F	١.
	NT. 0				

110. 4		
Whole Soya Flour	25	lb.
Whole Wheat Flour		lb.
Clear	50	lb.
Dry Milk	3	lb.
Salt	1.75	lb.
Shortening	2	lb.
Yeast	2	lb.

Sugar	1.5	lb.
Dry Malt	1.5	lb.
Water	about 10	gal.

The straight dough method is employed. A rather wide range in the quantity of water to be used is permitted. This is done to allow for the particular water absorption of the whole wheat flour and the clear that may be used by the baker. A straight dough is made but the whole soya flour is soaked for half an hour with a portion of the water before the dough is made.

"Non"-Staling Bread U. S. Patent 2,009,440

One-half to one per cent arabinose (based on flour) is added to dough.

Infant's Cereal British Patent 416,149

Wheatmeal	52.5 lb.
Oatmeal	18 lb.
Cornmeal	10 lb.
Wheat Germ	15 lb.
Salt	0.5 lb.
Lucerne	1 lb.
Dried Yeast	1 lb.
Bone Meal, Edible	2 lb.

100 lb. of above mixture is cooked with 35 gal. water and then dried on a heated drum.

Storage of Grain and Cereals, Improved British Patent 429,920

1000 lb. of solid carbon dioxide is used per 214 long tons of grain. Both are fed in simultaneously when loading ships or silos.

Chocolate Fudge

Unsweetened Chocolate	6	oz.
Sugar	2	lb.
Milk	1	lb.
Dairy Butter	1/3	oz.
Vanilla Extract	1/16	oz.

Cook slowly the melted chocolate, sugar, milk, and butter mixture to approximately 235° F. until a soft ball is formed when dropped into water. Remove from fire, add vanilla, beat thoroughly until the mass thickens, and then pour into well-buttered tin.

Chocolate Cream Fudge

Sugar	11/2	lb.
Corn Syrup	2	oz.

Unsweetened Chocolate	3	oz.	
Salt	1/16	oz.	
Evaporated Milk	8	oz.	

Heat to a boil, approximately 240° F., the mixed ingredients, until a soft ball is formed when dropped in cold water. Cool to approximately 100° F., and beat to a creamy consistency.

French Candy Balls

Unsweetened Chocolate	16 oz.
Powdered Sugar	2 oz.
Condensed Milk	16 oz.
Chocolate Topping	2 oz.

Melt chocolate in double boiler, add sugar and stir to prevent lumps. Add condensed milk and stir until smooth. Let set in cool place for two hours. Roll mixture into balls of desired size, and then roll balls in plate of chocolate topping. Let stand over night.

Jellied Fruit Candies

Plum Pulp	20 lb.
Peach Pulp	20 lb.
Cane Sugar	22 lb.
Corn Syrup	20 lb.
Powdered Pectin	1 lb.
Water	2 gal.

The pectin is mixed well with 5 lb. of sugar. This mixture is then stirred into the two gal. of water. Cook this solution slowly, to almost the boiling point, with stirring. Then to this smooth solution add the other ingredients. The entire batch is now cooked to 223° F. with stirring. The hot batch may now be deposited in starch molds, and allowed to become cold and firm.

Jellied Orange Candy

Pulp from 50 Oranges.		
Cane Sugar	35	lb.
Corn Syrup	25	lb.
Powdered Pectin	22	oz.
Citric Acid	6	oz.

The pulp is prepared by chopping up the oranges, and then cooked with twice its volume of water until soft, and then rubbed through a coarse strainer, to remove the seeds.

The powdered pectin should be previously mixed with about 10 lb. of sugar. The batch is now cooked to 223° F. Now dissolve the citric acid in a pint of water, add to the batch and once more cook to 223° F. The hot batch is now deposited in starch molds. Allow to become cold and firm.

Jellied Grape Juice Candy

Concord Grape Juice	3 gal. 18 lb.
Cane Sugar	18 lb.
Glucose or Invert Syrup	18 lb.
Powdered Pectin	13 oz.

The pectin is mixed well with about 5 lb. of cane sugar. This mixture is then stirred slowly into the grape juice. The batch is now slowly brought to a boil and then the balance of ingredients are added. Cook to 223° F. with stirring. The hot batch is now run into molds and allowed to cool.

Jellied Apple Juice Candy

Apple Juice (from Cooked	
Apples)	3 gal.
Cane Sugar	18 lb.
Glucose or Invert Syrup	18 lb.
Powdered Pectin	10 oz.
Citric Acid	5 oz.

Proceed as directed under Jellied Grape Juice Candy.

Jellied Pineapple Juice Candy
This juice can be used in the same
manner as outlined for grape juice.

Candied Sliced Orange, Lemon, and Grapefruit Peels

Slice peel about ¼ in. wide and 3 in. long. Cook peel with several changes of water to remove the bitterness and to make the peel tender. Now add to the peel about a 40% solution of sugar syrup (about 3 lb. sugar to the gallon of water) and cook until the temperature on the thermometer registers 217° F. Now drain the peel and allow to dry over night. The peel may be rolled in granulated sugar if desired. The peel can also be colored red or green with certified food color, if desired. Do this when cooking the peel in the last wash water.

Ginger, Preserved

Drain the ginger well and then cut it up. Place in cold water in a steam-pan, gently bringing to the boil and simmering for twenty minutes. Place in sieves to drain. Transfer to a cold syrup of 4 lb. sugar to each gallon of water, and allow to stand until next day. Transfer all to steam-pan, gently bring to the boil and simmer for 15 minutes. Then place in a clean dry tub and allow to stand until next day. Run off the syrup into the steam-pan and add 3 lb. sugar to each gallon of syrup. Stir well and

bring to the boil. Return this syrup to the ginger in the tub and allow to stand until the following day, then placing in sieves to dry. Roll in sugar and shake out the loose sugar through a coarse sieve. Lastly, spread out to dry.

Preserving Fruit Peels U. S. Patent 1,980,013

A process for treating the rind and peel of citrus fruits comprises heating the material in a sugar syrup for a period not to exceed about 1 hour at a temperature from about 212° F. to about 220° F., placing the material in containers with a relatively small quantity of sugar syrup, heating the containers, while they are unsealed, for a period of about half an hour at a temperature from about 212° F. to about 240° F, sealing the containers, and heating them for a period of about half an hour at a temperature of about 212° F. to about 240° F.

Preserving Red Raspberries by Freezing
The best result is obtained by freezing
at —18° F. in 50% syrup in either airtight or non-airtight containers, and then
storing at —12° F.

Pickling Vinegar Essence

The following is a formula for a concentrated liquid for making pickling vinegar:

Oil of Pimento	1/2	fl. oz.
Oil of Nutmeg	30	minims
Oil of Clove	90	minims
Tincture of Capsicum	1/2	fl. oz.
Acetic Acid (B.P.)	20	fl. oz.

One teaspoonful of this essence is mixed with each quart of vinegar to spice it.

Chewing Gum Bases

a. Bubble Gum Base:

• == ==================================	Ordina Dices		
Washed	Pontianac Gum	425	lb.
Washed	Gutta Katian	400	lb.
Washed	Gutta Soh	75	lb.
Candelil	la Wax	10	lb.

The mixed gums and wax are heated until the total batch contains only 8-9% moisture.

b. Stick Gum Base:

Pontianac Gum	425	lb.
Gutta Katian	400	lb.
Gutta Soh	75	lb.
Candelilla Wax	60	lb.
	Gutta Katian Gutta Soh	Gutta Katian 400 Gutta Soh 75

Chewing Gum Formula No. 1

2 011110100 2100 2		
Ball Gum:		
Base b (above)	22	lb.
Corn Syrup	48	lb.
Sugar	117	lb.
Chicle	3	lb.
Wax	$1\frac{1}{2}$	1b.
Caramel Paste	$2\frac{1}{2}$	
Flavor	$2\frac{2}{3}$	oz.
No. 2		
Penny Stick Gum:		
Base a (above)	40	1b.
Corn Syrup	40	lb.
Sugar	140	lb.
Flavor	30	oz.
No. 3		
Bubble Gum:		
Base a (above)	35	lb.
Pontianac Gum	5	lb.
Corn Syrup	45	lb.
Sugar	115	lb.
Flavor	28	oz.

Maraschino Type Cherries

Lambert, Royal Anne, Black Republican and Waterhouse varieties can be The fresh fruit should show a content of soluble solids in the juice of 16-18% at 21° C. and should be underripe rather than overripe. The bleach solution consists of sulphur dioxide (1.5%) together with sufficient air-slaked lime (5.4 lb. per 100 gal. of bleach) to keep the fruit firm and turgid. The cherries lose 7% in weight during the bleaching process. Approximately 250 lb. of cherries is stored in standard 52 gal. barrels and the strength of the bleach solution checked every few days by titration with standard 0.1 N 1 solution. Following bleaching, the cherries are stemmed, graded and pitted. sulphur dioxide-lime solution is removed by leaching with hot and then with cold water. The sulphur dioxide remaining in the cherries should be less than 0.035%. The dye used for coloring the cherries is No. 80 Ponceau 3R. A solution of % oz. of dye powder in 8 gal. of water is sufficient to color 100 lb. of pitted cherries at a temperature of 93° C. After coloring, the cherries are preserved by gradually increasing the concentration of sugar until a 50% syrup is reached. For flavoring, oil of bitter almonds and amyl acetate are used as desired. Pasteurization of the bottled cherries is effected by a heat treatment of 35 minutes at 91° C. for No. 10 cans holding somewhat less than 1 gal.

Preventing Browning of Peaches After Lye Peeling

Dip in ¼% hydrochloric acid for ½ to 1 minute and wash with water.

Non-"Bleeding" Jellies U. S. Patent 1,913,576

To prevent watering of jellies made with peetin, or agar, use ½ to 1% sodium alginate.

Jam and Jelly from Fruit Juices

Although most fruits contain small quantities of peetin and acid, many fruits do not contain sufficient amounts of these essential elements to produce jellies when the fruit juices are cooked with sugar. A small quantity of malic acid is found in the apple, and a little tartaric acid in the grape. Citric acid is contained in the lemon, the orange, and many other fruits.

Manufacturers of jellies can make high grade pure fruit jelly from all fruit juices by adding a very small amount of fruit acid (either citric, tartaric, or malie), less than one-half of 1%. The addition of small quantities of fruit acid and fruit pectin to fruits which are naturally deficient in these important constituents will improve the fluit flavor in the finished fancy preserve and the standard jam.

There are a few fruits which naturally contain enough acid and pectin to make jellies when the boiling with sugar is continued for 15 or 20 minutes. This excessive boiling, however, evaporates a large quantity of the fruit juice and flavor which should be retained in the finished product. For making jellies from these fruits deficient in pectin and acid, additional quantities of these substances must be added.

Purified powdered pectin is now being made from apples, lemons, and oranges by several firms. The product is very carefully standardized on the basis of jell strength, so that ½ oz. of purified powdered pectin will jell 50 oz. of cane sugar when mixed thoroughly with the sugar and then placed in a suitable cooking pan containing 2½ pints of water. Heat with constant stirring over a strong flame until the mixture weighs exactly 5 lb., then add ½ of a fluid oz. of a 50% solution of fruit acid. Mix thoroughly and pour into jelly glasses. Purified powdered pectin of such strength is designated 'No. 100.'

Pectin syrup is made by mixing thoroughly 5 lb. of powdered No. 100 pectin

with 20 lb. of cane sugar. Place the mixture in a suitable container and add sufficient boiling water to make 10 gal. when the temperature of the syrup is reduced to 70° F. Agitate a few minutes to dissolve the pectin. A 50% solution of fruit acid is made by placing 20 lb. of crystal, granular, or powdered tartaric, or citric acid in a 5-gal. stone jar and adding sufficient boiling water to make 5 gal. when the temperature of the liquid is reduced to 70° F. Agitate the hot liquid until the tartari, acid is dissolved.

All fruit juices for jelly production should have as little added water as is consistent with the proper extraction of pectin, color, and flavor from the fruit being used. Soft juice fruits, such as grapes, require very little, if any, additional moisture. Hard fibrous fruits, such as quinces, require the addition of a relatively large amount of water. In the following formulas for jellies, actual fruit juice is specified exclusive of added water. If water is added to the fruit during cooking, the amount of juice used in the formula should be increased by an amount equal to the quantity of water added at the time the fruit was heated in preparing it for the press, less the small quartity lost in evaporation.

Loganberry, Guava, or Pomegranate Juice Jelly

Filtered Fruit Juice (About 12 gal.) 100 lb. Cane Sugar 97 lb.

Fruit Juice from 2x1 Cold Pack Fruit (About 17 gal.) 167 lb. 2x1 cold pack fruit means 2 parts of fruit and 1 part sugar, usually placed in barrels and frozen.

Cane Sugar 30 lb. Fruit Pectin Syrup 1½ gal. 50% Solution Fruit Acid 10 fl. oz.

Crab-Apple, Currant, Gooseberry, or Quince Juice Jelly

Filtered Fruit Juice
(About 12 gal.) 100 lb.
Cane Sugar 97½ lb.
Fruit Pectin Syrup 1¼ gal.
50% Solution Fruit Acid 12 fl. oz.

Cherry, Elderberry, Strawberry, Pineapple, or Raspberry Juice Jelly Filtered Fruit Juice (About 12 gal.) 100 lb. Cane Sugar 96 lb.

or

Fruit Juice from 2x1 Cold
Pack Fruit (About 17
gal.) 167 lb.
Sugar 29 lb.
Fruit Pectin Syrup 2 lb.
50% Fruit Acid Solution 20 fl. oz.

Blackberry, Grape, or Plum Juice Jelly Filtered Fruit Juice (About 12 gal.) 100 lb.

Cane Sugar 97 lb.
or
Filtered Fruit Juice from
2x1 Cold Pack Fruit (About

17 gal.) 167 lb. Cane Sugar 30 lb. 50% Fruit Acid Solution 15 fl. oz.

Apricot, Peach, or Nectarine Juice Jelly Filtered Fruit Juice

(About 12 gal.) 100 lb. Cane Sugar 96 lb. Fruit Pectin Syrup 2 gal. 50% Fruit Acid Solution 23½ fl. oz.

In each formula for fruit jelly, cook the fruit juice, sugar, and fruit pectin syrup to 220° F. at or near sea level, or 8° above the boiling point of water in your factory. Then add the fruit acid solution and mix thoroughly. Fill the jelly quickly into glass and seal at once. If the temperature falls below 18° F. when the container is closed, it should be pasteurized at 180° F. for 20 minutes, if the glass does not contain more than 8 oz. The yield is approximately 164 lb. of finished jelly at 65% soluble solids.

Standard Cherry Preserves and Jam Fruit 82 lb. Cane Sugar 96 lb. Fruit Pectin Syrup 2 gal.

Fruit Acid Solution (50%) 13½ fl. oz.

In making fancy and standard preserves and jams, cook the fruit, sugar and pectin syrup to 221° F. at or near sea level, or 9° above the boiling point of water at your factory. Then add the fruit acid solution and mix thoroughly. The yield is approximately 158 lb. at 68% soluble solids.

In making standard preserves and jams, cook the fruit, sugar and fruit pectin syrup to 222° F. at or near sea level, or 10° F. above the boiling point of water at your factory. Then add the fruit acid solution and mix thoroughly. The yield is approximately 158 lb. at 68% soluble solids. Fancy preserves, jams, and standard preserves and jams

should pass through a cooling pan to reduce the temperature to 180° F. and then run quickly into glass, and be sealed at once. Then pasteurize glass containing from 1 to 2½ lb. at 190° F. for 25 minutes. The temperature is reduced to 180° F. before running preserves into glass so as to prevent the fruit from floating.

The thermometer should be accurate and should be tested at least once weekly when used daily. A very accurate determination for soluble solids contained in fruit products can be obtained by

using a refractometer.

Quince, Damson Plum, Gooseberry or Loganberry Jam

Fruit	100	1b.
Cane Sugar	$98\frac{1}{2}$	1b.
or		
Cold Pack Fruit	150	1b.
Cane Sugar	$48\frac{1}{2}$	1b.
Fruit Pectin Syrup	3	qt.
Fruit Acid Solution		-
(50%)	$7\frac{1}{2}$	fl. oz.
, , , ,		

Blackberry, Grape, Strawberry, Raspberry, Pineapple, or Plum Jam (Except Damson Plum)

Fruit 100 lb.
Cane Sugar 07

Cold Pack Fruit, 2x1 150 lb.
Cane Sugar 48 lb.
Fruit Pectin Syrup 1 gal.
Fruit Acid Solution (50%) 14 4. oz.

Apricot, Peach, Nectarine or Pear Jam
Fruit 100 lb.
Cane Sugar 97½ lb.
Fruit Pectin Syrup 1½ gal.
Fruit Acid Solution 18½ fl. oz.

 $\begin{array}{cccc} \text{Cherry Preserves and Jam} \\ \text{Fruit} & 100 \text{ lb.} \\ \text{Cane Sugar} & 96 \text{ lb.} \\ \text{Fruit Pectin Syrup} & 2 \text{ gal.} \\ \text{Fruit Acid Solution (50\%) 14 fl. oz.} \end{array}$

Damson Plum, Gooseberry or Loganberry
Jam

O 00		
Fruit	82	lb.
Cane Sugar	100	lb.
or		
Cold Pack Fruit, 2x1	123	lb.
Cane Sugar	571/2	1b.
Fruit Pectin Syrup	3	qt.
Fruit Acid Solution		- 7
(50%)	7	fl. oz

Blackberry, Grape, Strawberry, Pineapple, Raspberry or Plum Jam (Except Damson Plum)

Fruit	82	lb.
Cane Sugar	98	lb.
or		
Cold Pack Fruit, 2x1	123	lb.
Cane Sugar	$57\frac{1}{2}$	lb.
Fruit Pectin Syrup	1	gal.
Fruit Acid Solution		
(50%)	$13\frac{1}{2}$	fl. oz.
. , ,		

Apricot, Peach, Nectarine or Pear Jam
Fruit 82 lb.
Sugar 97½ lb.
Fruit Pectin Syrup 1½ lb.
Fruit Acid Solution (50%) 13½ fl. oz.

Cherry Pie Filler

Red Sour Pitted Cherries	450	lb.
Granulated Sugar	135	lb.
Corn Starch	25	lb.
Tapioca Flour	5	lb.
Water	9	gal.
Syrup	7	gal.
Benzoate of Soda	9	oz.

Put the cherries, sugar, syrup and benzoate into a steam kettle with 7 gal. of water. Bring to the boiling point and then add slowly while stirring, the paste made by mixing the corn starch and tapioca flour with the other 2 gal. of water. Heat and stir until the requisite consistency is attained.

Honey Jelly

Strained Honey	24	lb.
Citrus Pectin No. 80	4	oz.
Citric Acid Solution (50%)	1	oz.
Water	1	gal

Heat the honey to 155° F. in a steam-jacketed kettle.

In another kettle, heat the water to 180° F. Take about a pint of the honey and mix it with the dry pectin to make a smooth paste. Scrape this paste carefully into the hot water and bring to the boiling point, stirring until all is dissolved.

Add this solution to the honey and mix well. The resulting temperature should be 170° F. If not, raise to this point. Stir in the acid at once and run into containers.

For large size containers, 30 lb. pails or larger, use 20% more pectin.

Plum Jam

27	lb.
$12\frac{1}{2}$	lb.
50	lb.
4	oz.
$1\frac{1}{4}$	oz.
	12½ 50 4

Sugarless Marmalade for Diabetics

Lemons 11/2, the peel of one large orange, saccharin 5 gr., water 7 oz., gelatin 1/4 oz. Wash the orange and lemons, finely shave the skin (avoiding white pith) and chop up small; add the juice and pulp of the lemons. Put into saucepan and cover with the water. Bring to boiling point and simmer for two hours, adding water when necessary to keep to stated amount. Cut the gelatin into fine strips; add it with the saccharin to the mixture and stir for ten minutes. Put it into a jar an leave it to set. The keeping properties of this marmalade are not very good, and if it be desired to store it for any length of time a small quantity of sulphurous acid-forty parts per million—preferably in form of potassium metabisulphite should be added.

Apple Chutney

Apple chutney is prepared from the fresh apples, peeled, cored, and cut into pieces about half an inch cube. The exact shape of the pieces is not important so long as they are not too small. The apples, after chopping, are allowed to stand over night and then drained from any juice that may have separated,

the latter being reserved.

To every 60 lb. of apples 100 lb. of sugar is weighed out, made into a syrup with water, and boiled to 240° F. Into this syrup the small quantity of juice that may have separated is incorporated. While still boiling hot the syrup is poured on to the chopped apples in a suitable container, stirred and allowed to stand for 24 hours. The syrup and apple is then placed in a pan and boiled gently, together with chopped raisins, chopped stem ginger, and as much spice (such as mace, pimento, and nutmeg) and vinegar as taste demands, and the product bottled hot. Served with cold meatparticularly ham and pork-and similar dishes, this chutney is delightful. color should be golden brown, but this can be darkened if desired with a little sugar caramel. The only machinery required, apart from the boiling pan, is a chopping or dicing machine.

Apple Sauce

Apple sauce, well known in every home as the correct adjunct for roast pork and duck, and usually consisting of apples sliced and stewed with a little sugar, can be truly called a sauce if prepared as follows:

Fresh apples, as green and fresh as can be obtained, are placed in a clean barrel. A steam coil is inserted and the apples cooked for 15 minutes by contact with live steam at about 60 lb. pressure. Care should be taken to see that the steam line is drained before the valve is opened, otherwise the condensed water will enter the barrel and materially affect the consistency of the finished product.

When cooked, the apples are passed through a pulping machine, using the

finest sieve obtainable.

To 80 lb. of this apple pulp in a boiling pan add 80 lb. of sugar and 5½ lb. of 80% acetic acid. Stir and cook for 15 minutes. Spices (such as cinnamon, cloves, mace, and a trace of onion or garlic) may be added to suit individual taste, and the product filled into widemouthed bottles.

Apple Butter

Apple butter, which enjoys considerable popularity, is a preparation of a different type, being intended as a spread for sandwiches and at the teatable, and being in fact a kind of concentrated jam.

Processes vary, but consist in the main in expressing the juice from 100 lb. of freshly cooked apples and concentrating with 70 lb. of sugar in a boiling pan to 234° F. At this point 50 lb. of apple pulp, prepared as in the foregoing formula for apple sauce, are added, together with cinnamon, clove and mace spicing, and the mass gently cooked to 228° F.

Prevention of Browning of Fruit and

Treatment with a 0.1% solution of thiourea prevents or retards browning of surfaces of cut fruits. Addition of 0.01% thiourea to apple juice prevents darkening.

Chevon Mince Meats

Chopped Chevon (Cooked		
before Chopping)	10	1b.
Brown Sugar	15	lb.
Washed Currants	15	lb.
Molasses	10	lb.
Granulated Sugar	10	lb.

Seedless Raisins	15	lb.	
Chopped Apples	30	lb.	
Vinegar, Grape Juice, or			
Sauterne Wine	7	lb.	
Strong Coffee (Percolated			
Preferred)	10	lb.	
Jelly (Apple, Currant, Ras	sp-		
berry, or a Mixture)	5	lb.	
Citron	5	lb.	
Salt	1/2	lb.	
Lemons (Use Juice and			
Grated Rind Only)	$1\frac{1}{2}$	doz.	

Cook slowly for 3 hours, adding sufficient water to prevent burning. When cool, add

Rose Water	4 oz.
Cloves	4 oz.
Cinnamon	8 oz.
Nutmeg	4 oz.

Chevon from 8 months to a year old is best for this formula. In using this formula in pies place butter freely over surface before placing top crust.

Salted Soya Beans

A product similar to salted peanuts is obtained as follows:

Soak beans in salted water for 18 hours. Cook beans in lard or vegetable shortening at 170° C. until all water has been driven off and the beans float in the oil.

Fruit Gelatin Powder

Sucrose	30	lb.
Dextrose (Corn Sugar)	30	lb.
Gelatin (175 Bloom) 1.	5 - 2	lb.
	75-1	
Fruit Juice, Fruit and Water	37	lb.
Flavor and Color as	desi	$_{ m red}$

The gelatin is mixed in the water and dissolved in the usual manner, the sugars are dissolved and at 145° F. mixed with the gelatin solution. Cool to 100° F. and add remaining liquids such as flavoring color and acid. Let mixture thicken before adding fruits.

Pour into shallow pans to a depth of ¼ to ½ in. and set in cooler. When set turn out and cut into squares.

About 15% by weight of these cubes are stirred into ice cream as it comes from the freezer. The cubes may be added to the ice cream just before withdrawing but some naturally will be broken up.

A slab of the gelatin can be used as a layer in parfait ice cream and the cubes can be used as fillers in fancy pies, etc.

Jelly "Crystals"

Formula No. 1	
Sugar	90 lb.
Gelatin	20 lb.
Tartaric Acid	32 oz.
Flavor	6-8 oz.
Color	as desired
No. 2	
Sugar	31 lb.
Powdered Gelatin	5 lb.
Tartaric Acid	4 oz.
Flavor	2 oz.
Color	as desired

Gelatinless Jelly Powder U. S. Patent 1,974,474

Agar-Agar, Powdered	1/16	oz.
Sugar	$1\frac{1}{16}$	oz.
Tartaric Acid	1/16	oz.
Sodium Bicarbonate		gr.
The above forms a stiff	ielly with	8 oz.

The above forms a stiff jelly with 8 oz water.

Lemon Gelatin Powder

Sugar	10	lb.
Gelatin	1	1b.
Citric Acid	2.8	OZ.
Lemon Oil, U.S.P.	11/2	dr.
Certified Yellow Food		
Color	6	gr.
Water	61/	gr. fl. dr.

Blancmange Powder

Cornflour	100 lb	١,
Arrowroot	12 lb	١.
Color	12 dr	٠,
Flavor	6 oz	٠.

Custard Powder

Cornflour (St. Vincent)	300	lb.
Arrowroot	20-30	lb.
Vanilla	6	oz.
Essence Nutmeg	11/2	dr.
Color Powder	35 -	dr.

This to be used at the rate of 1½ oz. per pint of milk. The smoothness of the product is increased by the amount of cornflour used.

Compound Maple Table Syrups

Cane	Sugar-Map	le Sugar	Blends	
	Syrup		85 pt.	
Vermo	nt Maple Syr	rup	15 pt.	
Corr	Syrup—Can	e Sugar	Blend	
Corn S	Syrup (39° B	é.)	50 pt.	
	Syrup		50 pt.	

Caramel color to suit.

Invert Syrup	50 pt.
Sugar Syrup	50 pt.
Caramel color to suit.	
Cane Sugar—Molasses	Blend
Sugar Syrup	50 pt.

Cane Sugar-Invert Syrup Blend

New Orleans Molasses 50 pt.

Sugar Cane Table S	Syrup	
Sugar	7	lb.
Lemon Juice	3	oz.
Cream of Tartar	2	g.
Caramel Color	4	g.
Sugar Cane Syrup	5	οz.
Water	41/2	pt.
Benzoate of Soda	1/4	oz.

Dissolve the sugar in boiling water, then stir in the lemon juice and cream of tartar and color; then add the syrup and benzoate of soda. Boil for a few minutes and strain through fine muslin.

Chocolate Sauce

Unsweetened Chocolate	2	oz.
Dairy Butter	3/4	OZ.
Water	4	oz.
Sugar	1	lb.
Vanilla Extract	1,4	oz.

Melt the chocolate, and add the butter, stir until thoroughly mixed. Then add boiling water gradually with constant stirring. Heat to 230° F. and discontinue heating when a small portion cooled on a dish shows the proper consistency. Cool to approximately 100° F., and add the vanilla flavoring, stir thoroughly. This sauce can be used hot or cold.

Apricot Flavor

Linalyl Format	e ·	11/2	oz.
Amyl Valeriana	te	1/2	oz.
Oenanthic Ether	r	3/4	OZ.
Aldehyde C ₁₄		1/8	OZ.
Benzaldehyde		1/4	oz.
Peach Flavor		8	oz.
Glycerin		1	pt.
Alcohol		67	oz.
Water		34	oz.

Banana Flavor

Amyl Acetate		3	OZ.
Butyl Butyrate		1/8	OZ.
Isobutyl Ketone		1/4	oz.
Ethyl Benzoate		1/8	oz.
Orange Oil		1/4	oz.
Benzyl Valerianate			oz.
Cinnamon Oil, Ceyl	lon	15	min.
Mace Oil		30	min.
Heliotropin		1/4	oz.

Glycerin	52	oz.
Water	5	pt.
Alcohol	3	pt.

Burnt Almond Flavor

Caramel Color	2	oz.
Glycerin, C.P.	2	oz.
Benzaldehyde	$\frac{1}{2}$	oz.
Alcohol	8	oz.
Water	31/2	oz.

Cream Soda Flavor

Vanillin	5	oz.
Coumarin	3	oz.
Alcohol or Glycopon S	1/2	gal.
Glycerin		gal.
Water	1/4	gal.
One ounce will flavor five	galle	ons.

Kola Beverage Flavor

Grain Alcohol	$5\frac{1}{2}$	gal.
Best Vanilla Extract	14	oz.
Oil of Lemon	14	OZ.
Oil of Sweet Orange	7	oz.
Oil of Cassia	21	fl. dr.
Oil of Limes	4	oz.
Oil of Nutmeg	10	fl. dr.
Oil of Neroli	3	fl. dr.
Extract of Coca Leaves	1	fl. dr.

Allow to stand a month or more and then filter.

Maple Flavor Formula No. 1

roimua no. 1	•
Tincture of Foenugreek	6 pt.
Vanillin	3/4 OZ.
Musk	½ oz,
Balsam Peru	1 oz.
Oil Chamomile	⅓ dr.
Oil Celery	⅓ dr.
Tincture of Coffee	2 pt.
No. 2	

Foenugreek Oleoresin 5 lb. Hot Water 3 gal. Alcohol 1 pt. 15 oz. Malic Acid Compound Vanilla Extract 10 oz. Caramel Color 5 pt.

Rve Bread Flavor

Simple Syrup

150 oz.

Cumin Seed, Ground	11	lb.
Anise Seed, Ground	22	lb.
Coriander Seed, Ground	22	lb.
Caraway Seed, Ground	45	lb.

If a liquid flavor is desired the above is percolated with alcohol or if a nonalcoholic flavor is wanted Glycopon S is used.

	710, DEVELLAGED, PLAYONS 10
"Cloudy" Orange Syrup Concent	Household Extracts
Gum Arabic 24 Oil Orange Californian 34 Oil Lemon Californian 1 Orange Color Solution 18 Simple Syrup 72 Sulphonated Castor Oil 4 Weter to make 128	oz. (Alcoholic)
Oil Lemon Californian 34	Pure Lemon Extract
Orange Color Solution 18	Lemon Oil 6.4 oz.
Simple Syrup 79	Alcohol, Pure 115 oz.
Sulphonated Castor Oil 4	oz. Lemon Oil 6.4 oz. oz. Alcohol, Pure 115 oz. oz. water to 1 gal.
Water to make 125	Pure Orange Extract
Pass through colloid mill.	Orange Oil 6.4 oz. Alcohol, Pure 115 oz. Water to 1 gal.
MACHINE STREET, AND CONTROL OF STREET, AND CO	Alcohol, Pure 115 oz.
Dwied Plackbowy Concentrate	Water to 1 gal.
Dried Blackberry Concentrate	Down Almond Tretmost
	lb. Pure Almond Extract
	pt. Oil Bitter Almond,
Water 4	pt. F.F.P.A. 1.28 oz.
Martine a seguina e con de contra en el cont	pt. pt. Coil Bitter Almond, pt. F.F.P.A. 1.28 oz. Alcohol, Pure 40 oz. Water to 1 gal.
Cherry Concentrate, Natural	
Cherries, Dried 8	lb. Imitation Vanilla Extract
	pt. Vanillin 70 oz.
Water 4	pt. Coumarin 4/4 oz.
Put cherries in water, heat, cool	and Alcohol, Pure (25% by
add alcohol.	Volume) 25 gal.
	Simple Syrup 80 oz.
Q	Vanillin 70 oz.
Cognac Essence	Imitation Lemon Extract
Cognac Ether 650 Rum Ether 650	g. Citral % oz. 6 g. Alcohol, Pure 5 pt. 6 g. Water to 1 gul. 6 g.
Rum Ether 650	g. Alachol Pura 5 nt
Sweetened "Saltpeter Spirit" 165	Water to 1 gal.
Ethyl Acetate 165	g. Water
Oenanthic Ether	2 g.
Sugar Color 555	Caraway Extract
Sweetened 'Saltpeter Spirit' 165 Ethyl Acetate Oenanthic Ether Sugar Color Alcohol (90%) 4000	Formula No. 1
The state of the s	Oil of Caraway 3 g.
Rum Essence	Oil of Caraway 3 g. Alcohol 50 g.
Rum Either 200	g. Glycerin 6 g.
Ethyl Acetate 40	Alcohol 50 g. Glycerin 6 g. 9 g. Water 41 g.
Cinnamon, Tincture 10	No. 2
Catechu, Tincture 10	g. Oil of Caraway 3 g. Alcohol 80 g.
Vanillin, Tincture 10	g. Alcohol 80 g.
Ethyl Formate 75	80 g. 80 g. 20 g. 20 g. 20 g.
Angelica Root, Tincture	2 g.
Peruvian Bark, Tincture 15	g.
Orange Flower Water 100	g. Cardamom Extract
Woodruff Essence 30	g. Oil of Cardamom, Ceylon 3 g.
Butyric Ether 20	g. Alcohol 50 g.
Alcohol (90%) 650	g. Glycerin 6 g.
Rum Ether 200 Ethyl Acetate 40 Cinnamon, Tincture 10 Catechu, Tincture 10 Vanillin, Tincture 75 Angelica Root, Tincture 2 Peruvian Bark, Tincture 15 Orange Flower Water 100 Woodruff Essence 30 Butyric Ether 20 Alcohol (90%) 650 Rum 1000	g. Water 41 g.
× value and the parameter of the paramet	1 '
Rock and Rye Whisky Essence	Cassia Extract g. Formula No. 1
Grain Fusel Oil Rectified 340 Green Wine Lees Oil 12 Peru Balsam 12	g. Formula No. 1
Green Wine Lees Oil 12	g. Oil of Cassia Rectified 3 g.
Peru Balsam 12	g. Oil of Cassia Rectified 3 g.
Jamaica Rum Essence 12	
Vanillin 6	g. Glycerin 6 g. Water 41 g.
Ethyl Acetate 12	g. Water
Coumarin 15	g. No. 2 (Cinnamon)
Raisin Wine Essence 580	g. 3% Standard
Peach Essence 8	g. Stanuaru
Bitter Orange Extract 50	g. Oil of Cassia Cinnamon 30 g.
	g. Alcohol 200 g.
Clove Oil 2.5	5 g. Water 170 g.

101	L CLLDIVII OIL	D I Oldin Oblitte i	
Extract Celery		Extract Juniper	
Formula No. 1		Oil of Juniper	2 g.
Celery Oil	0.6 g.	Alcohol	90 g.
Alcohol	600 g.	Water	8 g.
Water	400 g.	Banana Oil (Syntheti	(a)
manufacture of the second of t		lb. oz.	dr. min.
No. 2		Benzyl Acetate 2 15	7 32
Oil of Celery	0.5 g.	Benzyl Acetate 2 15	3 54
Alcohol	60 g.	Heliotropin – 1	2 - 58
Glycerin	6- g.	Vanillin - 1	2 58
Water	34 g.	Geranyl Acetate – –	$\begin{array}{ccc} 3 & 16 \\ 2 & 12 \end{array}$
		Geranyl Acetate – – Terpeneless Lemon	<u> </u>
Wild Cherry Extr		Oil	1 16
Wild Cherry Bark	8 lb.	Marin of the state	
Alcohol Water	4 lb. 4 lb.	Blackberry Oil	
Percolate and filter.	± 10.	Vanillin	2 g. 3 g.
rercolate and litter.		Coumarin	3 g.
Ciarana Tartar	.	Heliotropin Methyl Salicylate	2 g. 2 g. 1 g.
Cinnamon Extra		Methyl Anthranilate	1 g.
Oil of Cinnamon, Ceylon Alcohol	3 g. 50 g.	Orris (10% Solution)	5 g.
Glycerin	6 g.	Coriander Oil	6 g.
Water	41 g.	Fennel Seed Oil	18 g.
	G	Amyl Butyrate Ethyl Benzoate	112 g.
Clove Extract		Amyl Acetate	256 g. 192 g.
Formula No. 1		Ethyl Acetate	397 g.
Oil of Cloves	3 g.	Aldehyde C_{16}	4 g.
Alcohol	50 g.		
Glycerin	6 g.	Brandy Oil	
Water	41 g.	Green Cognac Oil	20 g.
		Oenanthic Ether	80 g.
No. 2	00	Rum Ether Fusel Oil	80 g. 20 g.
Clove Oil Alcohol	20 g. 650 g.		- S.
Water	350 g.	Oil Wild Cherry	
		Formula No. 1	
Coriander Extra	et	Benzoic Acid	4 g.
Formula No. 1		Benzaldehyde	6 g.
Oil of Coriander	3 g.	Amyl Butyrate Ethyl Acetate	6 g. 24 g.
Alcohol	50 g.	Ethyl Benzoate	$\frac{1}{24}$ g.
Glycerin	6 g.		_
Water	41 g.	No. 2	
Approximation and the second of the second o		Amyl Acetate	24 g.
No. 2		Amyl Butyrate Ethyl Benzoate	12 g. 12 g.
Oil of Coriander	3 g.	Benzaldehyde	32 g.
Alcohol	80 g.	Oil Sweet Orange Calif.	4 2.
Water	20 g.	Oil Cloves	3 g.
Ginger Ale Extra		Cherry Oil (Syntheti	
Oleoresin Capsicum	112 oz. 1 oz.	lb. oz.	dr. min.
Safrol Cinnamic Aldehyde	1 oz.	Benzylidene Formate 1 -	
Mace Oil	1½ oz.	mate 1 - Oenanthic Ether 4 8	
Citral	$1\frac{1}{2}$ oz.	Ethyl Methyl An-	
Geranyl Acetate	¹ / ₄ oz.	thranilate 1 6	3 12
Alcohol	1 pt.	Benzaldehyde,	1 10
One ounce will flavor five	gamons.	F.F.C. 3 1	4 48

Oil Cognac	Oil Pear Ethereal
× =	
Ethyl Butyrate 21 g.	Benzyl Propionate 1 oz. Amyl Acetate, Pure 11 oz. Butyric Ether, Absolute 4 oz.
Ethyl Butyrate 21 g. Oil Cognac 28 g.	Butyric Ether, Absolute 4 oz.
Tincture of Prunes 480 g. Ethyl Butyrate 21 g. Oil Cognac 28 g. Oenanthic Ether 42 g.	2009110 20101, 210001110
	"Scotch" Whisky Oil
Oil of Green Cognac	Fusel Oil Rectified 510 g. Cade Oil 84 g. Ethyl Butyrate 445 g. Bitter Almond Oil 20 g. Sweet Almond Oil 20 g. Guaiacum Oil 10 g.
Sebacic Ether 5 g.	Cade Oil 84 g.
Pelargonic Ether 2 g.	Ethyl Butyrate 445 g.
Pelargonic Ether 2 g. Cognac Oil 3 g.	Bitter Almond Oil 20 g.
Pelargonic Ether 2 g. Cognac Oil 3 g. Oenanthic Ether 90 g.	Sweet Almond Oil 20 g.
And a state of the	Guarastan on 10 g.
Cola Oil for Beverages Oil Lemon 120 g. Oil Sweet Orange 80 g. Oil Nutmeg 40 g. Oil Coinnamon 40 g. Oil Coriander 20 g. Oil Neroli, Artificial 40 g. Alcohol (75%) 15,360 g.	Oil Strawberry (Synthetic)
Oil Lemon 120 g.	oz. dr. min.
Oil Sweet Orange 80 g.	Ethyl Acetate 42 5 15
Oil Cirpower 40 g.	Aldehyde C ₁₆ 23 3 40
Oil Cariandar 20 g.	Amyl Acetate 12 6 24
Oil Noroli Artificial 40 c	Ethyl Butyrate 9 - 27
Alcohol (75%) 15 360 g	Amyl Butyrate 9 - 27
AROMOT (10/c) 10,000 g.	Propyl Iso Butyrate 58 5 15
C	Ethyl Formate 1 2 13
Curacao Oil	Oil Cognac, Green - 6 47
Curacao Oil	$ \begin{array}{cccccccccccccccccccccccccccccccccccc$
Oil Cassia 30 g.	Oil Raspberry (Artificial)
Timelyl Agetate 50 g.	
Potitore in Oil 75 o	
Oranga Oil 650 g	Tea Rose, Oil - 9 4 45 Aldehyde C ₁₆ - 11 5 25
Lemon Oil 150 g.	Amyl Cinnamic
2008	Formate 1 6 6 10
"Holland" Gin Oil	Vanillin
	Amyl Acetate - 10 3 25
Lemon Oil 3 g.	Ethyl Butyrate — 8 2 44 Ethyl Formate — 8 2 44 Ethyl Acetate — 12 4 6 Iso Butyl Acetate 2 11 6 20
Angelias Poet Oil 16 g	Ethyl Formate - 8 2 44
Final Oil Rectified 12 of	Ethyl Acetate - 12 4 6
Rosemary Oil 16 g.	Iso Butyl Acetate 2 11 6 20
Coriander Oil 13 g.	1so Cinnamic
Lemon Oil 3 g. Anise Oil 3 g. Angelica Root Oil 16 g. Fusel Oil Rectified 12 g. Rosemary Oil 16 g. Coriander Oil 13 g. Juniper Berry Oil 940 g.	Acetate 1 7 6 11
	Amyl Butyrate – 8 2 44
"Old Tom" Gin Oil	Concentrated Foam for Beverages
Coriander Oil 270 g. Anise Oil Rectified 80 g.	Saponin 16 oz. Glycerin 64 oz. Distilled Water 64 oz.
	Glycerin 64 oz.
Juniper Berry Oil Rectified 610 g.	Distilled Water 64 oz.
Caraway Oil 20 g. Angelica Root Oil 15 g.	Use 1 oz. to 15 gal. syrup.
Angenes not on 10 g.	G M: T G M
Oil Grape (Synthetic)	Caffein-Free Coffee
lb. oz. dr. min.	U. S. Patent 2,023,333
Oil Cognac Green - 14 5 26	Ground raw coffee is extracted with a warm mixture of
Methyl Anthran- ilate 7 2 3 55	aa-dichlorethane and
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	αβ—dichlorethane
Propyl Cinnamate - 5 6 58	
Ethyl Butyrate 1 1 4 55	Artificial Mineral Water
	Austrian Patent 142,032
Oil Kümmel Danzig	1 liter of following solution is mixed
Carvol 300 g.	with 10 liters of carbonated water:
Coriander Oil 3 g.	Salt 0.02 g.
Orange Oil 3 g.	Magnesium Sulphate 0.02 g.

Dihydrogen Sodium		Essence Orange	5 oz.
Phosphate	0.02 g.	Sulphurous Acid	4 oz.
Potassium Nitrate	0.008 g.	ourprise out and the	_ 02.
Calcium Oxide	0.2 g.		•
Calcium Oxide	o 8.	Tonic Water	
****		Quinine Bisulphate	8 gr.
Lime Barley Wat	er	Aerated Lemonade	4 pt.
	. 1	Aerated Water	4 pt.
Syrup (66°)	2 gal.		
Barley Extract	3 qt.	7 7 6	,
Refined Lime Juice	1 qt.	Lemonade Cryst	
Citric Acid Powder	7 oz.	Sugar	100 lb.
or	4.	Lemon Juice Powder	4 to 6 oz.
1-2 Solution	14 oz.	Tartaric Acid	4 lb.
Essence Lime	3 oz.	and any deployment of the second of the seco	
Sulphurous Acid	3 oz.	Orangeade Cryst	-ola
Lemon Color	½-1 oz.	Sugar	100 lb.
		Orange Juice Powder	
		Tartaric Acid	4 lb.
Orange Barley Wa	ter	Tartaric Acid	4 10.
Syrup (66°)	3 gal.	***************************************	•
Orange Concentrate 6-1		Lime Juice Crys	stals
Orange Beverage Base	7 pt.	Sugar	100 lb.
Barley Extract	1 gal.	Lime Juice Powder	2 to 4 oz.
Orange Color, if Desired	2-6 oz.	Tartaric Acid	4 lb.

SUGAR TABLE FOR SODA WATERS

Pounds of Sugar Added to 1 Gal. Water	Quantity of Syrup Obtained		Sugar Percentage in Syrup	Density	Degrees Baumé at 60° F.
1 2 3 4 5	gal. pt. 1 - 1 1 1 1 1 1 1 2 1 3	oz. 10 4 14 8 2	$10\frac{3}{4}$ $19\frac{1}{4}$ $26\frac{1}{2}$ $32\frac{3}{4}$ $37\frac{1}{2}$	1.043 1.080 1.113 1.142 1.166	6 11 15½ 18 20½
6	1 3	$ \begin{array}{r} 12 \\ 6 \\ \hline 10 \\ 4 \end{array} $	41¾	1.188	23
7	1 4		45¼	1.209	25
8	1 5		49	1.227	26¾
9	1 5		52	1.244	28¼
10	1 6		54½	1.258	29½
11	1 6 7 2 0 2 0 2 1	14	57	1.271	3034
12		8	59	1.284	32
13		2	51	1.296	33
14		12	6234	1.306	3334
15		6	644	1.315	341/2
16	2 2	$ \begin{array}{r} 10 \\ 4 \\ 14 \end{array} $	65½	1.324	34¾
17	2 2		67¼	1.332	35¼
18	2 3		68½	1.340	35%
19	2 3		69¾	1.347	36

Aging Alcoholic Liquors U. S. Patent 1,963,165

About a pound and a quarter of potassium permanganate crystals are dissolved in an appropriate amount of water, for example three and one-half gallons. To this solution there is added about a pound of sulphuric acid, pref-

erably concentrated sulphuric acid. The aqueous mixture resulting from mixing sulphuric acid and a water solution of potassium permanganate is added to the raw alcoholic liquor, preferably in the proportions of one pound of liquid to about fifty gallons of raw alcoholic liquor. The raw alcoholic liquor usually

comes in charred barrels provided with a removable bung. In operating according to the present process, the bung is removed from the barrel and the aqueous mixture resulting from mixing sulphuric acid and potassium permanganate in solution is added to the contents of the barrel. Thereafter the bung is replaced and the barrel and its contents are allowed to mature for a short period of time at an elevated temperature. Rye and bourbon are allowed to mature for about three days at a temperature of 120° F., while rum and brandy are allowed to mature for about two days at the same temperature. When using a lower temperature, for example 100° F., rye and bourbon can be allowed to mature for a period of five days, and rum and brandy for a period of four days. The important point is that after the treatment of the raw alcohol liquor with the treating solution there should be a short maturing period. The function of the elevated temperature is to accelerate the maturing period, and therefore, if the temperature is reduced, the maturing period at this point becomes longer and vice versa. If the temperature is increased above 120° F., the maturing period can be shortened. Of course, the upper temperature limit cannot be too high, since the treatment mixture at highly elevated temperatures would deteriorate the quality of the alcoholic

When the raw alcoholic liquor is treated with the aqueous solution resulting from mixing sulphuric acid and potassium permanganate, there is immediately set up in the liquor a substantial agitation, acting to eliminate the poisonous components of the fusel oils including the aldehydes and the higher alcohols while leaving the esters of the fusel oil to which the aromatic flavor of the liquor is due substantially unimpaired.

After the treated alcoholic liquor has been allowed to mature, as set forth above, the temporary bung is removed. When the bung is removed from the barrel, the chemical and physical action which the liquor is undergoing is very apparent. Immediately upon removal of the bung, there is an evolution of vapors and gases, these representing partial reaction products of the treatment process up to this point. A portion of the impurities present in the original raw liquor have been removed by virtue of the absorptive capacity of the porous lining of the barrel which, as stated, is also in a charred condition, thus augmenting the initial absorptive capacity

of the porous wood of which the barrel is made.

Immediately upon removing the temporary bung from the barrel there is added to the treated alcoholic liquor an agent which will function to bleach and stop the chemical and physical activity taking place in the liquor which has been treated with the sulphuric acid and the While various permanganate mixture. agents may be used to effect the bleaching and the cessation of chemical and physical activity in the alcoholic liquor, it has been found that most satisfactory results are obtained by the addition of an oxygen evolving agent. While the preferred oxidizing agent is hydrogen peroxide, other compounds which are the chemical equivalents of hydrogen peroxide may be used.

The amount of the bleaching and activity neutralizing agent which is added to the treated alcoholic liquor will of course vary with the character and quality of the initial raw product and with the amount of the sulphuric acid permanganate solution which has been initially added to the raw liquor. When adding the sulphuric acid permanganate treatment agent in the proportions above set forth to about 50 gal. of raw liquor, it has been found that the addition of eight ounces of 30% hydrogen peroxide

gives very satisfactory results. After the addition of the bleaching and activity neutralizing agent, a permanent bung is inserted into the barrel and the treated alcoholic liquor allowed to further mature, preferably under an elevated temperature. The following maturing procedure has been found to give most satisfactory results. When maturing rum and brandy, the barrels of alcoholic liquor treated in accordance with the previous steps of the process are maintained in a warehouse having a temperature of about 120° F. for about three weeks. Thereafter the temperature is reduced to about 100° for another week, and then to about 80° F. for an additional week. The warehouse or room in which the liquor is being matured under elevated temperature is then allowed to cool off to normal temperature which usually takes about a week or ten days. unless artificial means are used for cooling the temperature of the storage room. In general this period of maturity varies from about 6 to 8 weeks, and the resulting rum and brandy has reached full maturity, having a flavor and mellowness equivalent to rum and brandy which have been naturally aged for a period of approximately four years.

When rye and bourbon are treated, due to the higher content of impurities including fusel oils present in the raw alcoholic liquor, a longer period of maturing When rye and bourbon is necessary. have been treated as above set forth with the sulphuric acid potassium per-manganate solution, and then later on after a short period of maturing treated with the bleaching and activity neutralizing agent, the so treated material is subjected for a period of about two months to a temperature of about 100° F. The temperature of the storage room containing the barrels of treated liquor is then reduced to about 100° F. and the treated liquor allowed to mature for about an ad-Thereafter the ditional two months. temperature of the storage room is reduced to 80° F. for a period of one The storage room is then allowed to cool off to about 70° F., it taking about one month under average conditions for the storage room to reach this temperature, although it is recognized that the cooling may be accomplished much quicker by artificial cooling means.

Kümmel Danzig		
Carvol	300	g.
Coriander Oil	3	g.
Orange Oil	3	g.
Alcohol	5000	g.
Water	2250	g.
Glycerin	274	g.

Methyl, Isopropyl and Amyl Alcohols, Tests For

0.1 g. of vanillin is dissolved in 10 mils of alcohol in a test tube and 1 mil of pure sulphuric acid carefully run down the side of the test tube to form a layer at the bottom. By slightly rotating the tube the alcohol and acid are cautiously mixed (care is needed otherwise the sudden rise in temperature will cause violent ebullition) and the colors formed at the area of contact and of the final mixture are noticed.

Methyl Alcohol:	Area of contact Final mixture	Pale mauve Pale mauve
Ethyl Alcohol:	Area of contact Final mixture	Lemon yellow Colorless
Isopropyl Alcohol:	Area of contact	Bright red towards the acid layer changing to deep blue towards the alcohol layer
	Final mixture	Prussian blue
Amyl Alcohol:	Area of contact Dull red tow layer chan blue towar layer	
	Final mixture	Prussian blue A white precipitate also forms

In order to differentiate more accurately between isopropyl alcohol and amyl alcohol 10 mils of water are added to each mixture and then shaken well. With isopropyl alcohol the mixture becomes pale blue but rapidly fades, becoming water-white. With amyl alcohol the mixture separates into two layers, the upper alcoholic one being deep grass green (permanent after two hours) and the lower aqueous layer water-white. The white precipitate settles to the bottom.

Approximate Estimation

For the approximate estimation of methyl, isopropyl and amyl alcohols in ethyl alcohol the quantity of sulphuric acid used was increased to 3 mils. Dilutions of the three alcohols in ethyl alcohol were made, 1 in 10, 1 in 100, 1 in 1000 and 1 in 10,000, also a control of ethyl alcohol alone, the color obtained with the latter and the solution of vanillin being a distinct yellowish green.

	1 in 10	1 in 100	1 in 1000	1 in 10,000
Methyl Alcohol	Blue green	Very faintly blue green	Yellowish green	·
Isopropyl Alco- hol	Blue	Pale blue	Blue green	Yellowish green
Amyl Alcohol	Deep blue	Blue	Pale blue	Very faintly blue green

Preserving Brewer's Yeast Yeast will keep well indefinitely if covered by a 10% cane sugar solution.

Seed Yeast for Production of Commercial Yeast U. S. Patent 2,016,791

After mixing about 4 lb. yeast with an aqueous aerated "cream" formed by agitating about 12 oz. of calcium sulphate with water, 0.5 to 1.0% of corn starch is added to the mixture, it is maintained at a temperature of about 28° C. for about 30 hours, then diluted with aerated water and allowed to stand for about 18 hours to produce sporulated and durable yeast.

Isinglass Finings for Beer Clarification British Patent 432,159

Pieces of isinglass are steeped in acidified water for several hours and then gently stirred continuously for 12-15 hours. The liquid, which then has the consistency of thin treacle, is strained through a fine sieve.

Home Made Wine

To two volumes of water in a large glass bottle, add one volume of washed whole grapes and one volume of sugar. Stopper with a cotton plug, place in a warm place, shake up well daily, and allow to ferment for about 8 weeks or until the evolution of gases ceases. Then siphon off or decant, sweeten to taste, bottle and set aside to age.

Bee Wine

Four ounces of sugar and 4 oz. of treacle are mixed with 1½ pt. of water to form the mother liquid. Small pieces of the ginger beer plant are then added, and the mixture is kept in a warm place. Each day about a teaspoonful of sugar is added, there is brisk fermentation and a palatable drink is soon ready. The ferment quickly increases, and can be used to prepare a new batch.

Orange Wine

Cut well ripened oranges in half and squeeze out juice. Strain out coarse pulp and seeds. Add 150 p. p. m. of sulphur dioxide; corresponds to about 21/2 lb. of metabisulphite or about 14 lb. of sulphur dioxide per 1000 gal. of juice. Mix well. Add sugar to increase the Balling degree to 22-24° Balling for a dry wine of medium alcoholic content and to 32-33° Balling for one that will contain a small amount of sugar after fermentation is complete.

Ferment large quantities in open redwood vats, artificially cooling the fermenting liquid, if necessary, to maintain the temperature below 85° F. Smaller quantities are fermented in oak puncheons or barrels. Take the Balling degree once a day to follow the course

of fermentation.

fermentation ceases.

When fermentation becomes slow and is nearing completion transfer from the open vat to a covered redwood tank, leaving the bung hole open. Fit with a fermentation bung in order to give a slight pressure of carbon dioxide gas in the tank and thus prevent the growth of vinegar bacteria. Similarly equip barrels or puncheons.

When gas is no longer given off remove fermentation bung and fill the tank, puncheon or barrel with fermented orange juice and seal with an ordinary bung. Once or twice a week for several weeks loosen the bung for a few seconds to release accumulated gas pressure until

Then let stand for two or three weeks to settle, with bung tightly in place. Next drain off, that is, rack from the sediment; this can be done through a bronze spigot inserted in a bung hole near the bottom of the tank, or by syphoning by hose from smaller con-tainer. Transfer to clean cooperage that has been sulphured (in which a sulphur wick has been burned). Fill these containers completely full. Let settle two or three weeks. Then rack. Filter clear. This is easily done, usually by means of a pulp filter. The wine can then be polished brilliantly clear through a porcelain candle, or pad type polishing filter. It should next be aged, in wood as is done with grape wine. If new wood is used the tanks or barrels should be soaked out with dilute soda ash solution and water before use in order

a wood taste. If to be rapidly aged, heat to 120° F. a few days in the presence of about 1/2% by weight of oak chips if in redwood, and a head space of about 10%. If in oak barrels no chips are needed. Pump over occasionally. Do not overdo the rapid aging process. Watch carefully and stop the treatment when the desired amount of aging is attained. Try it first on a small scale, in order to avoid

that the wine will not acquire too strong

"grief" and loss by improperly rapid aging of a large quantity.

After aging, the "wine" may need a polishing filtering again. After filtering let it rest in wood a few days to

"recover" before bottling.

If a fortified wine is desired a special permit or license is required, numerous regulations must be met and numerous forms filled out, either to install a still and use brandy made on the premises for fortification, or to buy fortifying brandy of high proof. Having conformed to all regulations, etc., then brandy may be added to bring the wine to 20-21% alcohol. "Angelica" type sweet fortified orange "cider" should show about 10% sugar by chemical test and sherry type 2-4% sugar by chemical test. The former is aged like dry wine; the latter is heated at 130-140° F. for 2-3 months to acquire a sherry flavor and color. By gentle aeration the time can be greatly shortened.

"Champagne' type sparkling orange wine can be made by fermenting juice of 21-22° Balling dry; filtering; aging a few months; adding 2% of cane sugar; fermenting in the bottle with Champagne yeast, disgorging and refilling the bottles; or by fermenting in bulk by the Charmat or other bulk process; filtering and bottling under carbon dioxide

pressure.

Or the orange "wine," sweet or dry, can be carbonated with carbon dioxide gas in one of several types of carbonat-

ing machines.

In order that non-carbonated, nonfortified sweet "wine" after bottling will not undergo bacterial spoilage it may be preserved with about 300 p.p.m. of sulphur dioxide, or by pasteurizing in the bottle at 140° F. for 30 minutes.

Berry "Wines"

Here the procedure is somewhat different than in making orange "wine." Use ripe, sound berries, sorting out moldy fruit. Crush into open vats. Add 8 oz. of potassium metabisulphite or 4 oz. of sulphur dioxide per ton, or about 2½ lb. of the former or 1¼ of the latter to each 1000 gal. The metabisulphite is dissolved in water 8 oz. per gal. before addition. Add to the juice. Mix well. Wait 2 hours. Add a starter or 2–3% pure yeast culture. Stir or punch three times daily until the Balling degree drops to about ½ or ¼ the original Balling degree. Fermentation extracts the color and tannin and softens the fruit.

Press in a rack and cloth press. To the juice add for a dry "wine" 15% by weight of sugar; for a sweet "wine" about 25%; that is to 1000 gal. of the juice about 1350 and 2250 lb. of sugar respectively. See that it all dissolves.

Ferment and treat as described for

orange "wine."

Rhubarb Wine

Run 32 lb. rhubarb through a meat chopper, strain the juice into a vat and add 6 gal. water. Let stand for 2 days and strain. Let stand for 1 to 2 days, siphon off the clear liquid into a keg and add 24 lb. sugar. Boil up 2 lb. raisins in a little water and add together with 1 lb. sugar coloring. Add also a little gelatin as clearing agent. Let ferment for about 14 days, or until complete. Fill up keg with water and let stand for 4 months before tapping.

Dehydration of Fresh Soya-Slime German Patent 602,935 and 599,639

Example of a Soya-Mud of composition:

Water	50	oz.
Lecithin	40	oz.
Soya-Oil	10	θZ.

Warm

Soya-Slime 100 oz.

to 60° C., and add

Glycerin Containing Dry
Sugar (until sp. g. = 1.36
to 1.39)
25-50 oz.

Stir thoroughly ¼ hour, allow to stand. Two layers formed, the heavy one:

Glycerin + Water

+Sugar

the light one:

Lecithin + Oil +

Water

Repeat to get a water-content of 10%.

Defoamer for the Sugar Industry Prevents foaming when "saturating" the lime-containing "thin sap." Woolfat, Neutral.

For the Alcohol Industry:

Coconut Oil 80-85 Vaseline Oil 20-15

Preservation of Coffee U. S. Patent 1,956,290

Oxidation and "staling" of coffee is curtailed by addition of 0.3% sodium pyrosulphate.

Denaturation for Food Salt (per 100 kg.)

Formula No. 1

Mineral Oil No. 2

Iron Oxide O.25 kg.
No. 3

Soap Powder 1 kg.
For the Chemical Industry

| No. 4 | Sodium Sulphate, Crystallized | No. 5 | Sodium Carbonate | No. 6 | Crystal Ponceau 6R | O.5 | g.

Non-Caking Salt British Patent 407,829

The addition of up to 7% potassium chloride to granular table salt prevents caking.

Non-Caking Sugar

Caking of sugar is prevented by addition of 1% tricalcium phosphate.

Improving Liquid Honey

Heat honey to 71° C.; cool rapidly to 24° C.; add fine crystallized honey with stirring for 15 minutes; cool and bottle.

Non-Mottling and Non-Hardening Maple Sugar

U. S. Patent 1,970,870

Maple sap or syrup is boiled in an open vessel until the temperature reaches 125° F., then allowed to cool, and continuously stirred until cold. The crystallized mass obtained, containing about 2% of water is pressed into blocks occupying 30-31.4 cu. in. per lb.

Clarifying Cider

Pectin (20-30 oz.) is added to 1 gal. of warm eider and the mixture shaken

at intervals for 20 minutes. The strained liquor is added to 100 gal. of cider to be clarified, and after 15 hours at approximately 21° C. the cider is siphoned off, mixed with 2-3 lb. of diatomaceous earth, and filtered through canvas.

Wax Coating for Citrus Fruit U. S. Patent 1,940,530

Fresh fruit (notably citrus) is improved in appearance and made less liable to wither if a thin film of molten wax is rubbed on to the surface (e.g., 5-15% of carnauba wax in paraffin wax at 77-105° C. rubbed on for 10-30 seconds). Advantageous results are obtained if an alkaline wash has preceded this treatment.

Curing Ripe Olives U. S. Patent 1,928,229

Wash olives in ½ to 2% caustic soda solution then in water till neutral. Soak in ½ to 5% pyrogallol for a few hours. Without rinsing soak in 1% caustic soda solution until skin is penetrated; expose to air until black; wash till free from alkali and then soak in brine to develop flavor.

Storing Walnut Meats

Bleached nuts are preserved by packing in earthenware containers with alternate layers of coconut fiber and a 9 to 1 mixture of salt and sodium dihydrogen phosphate crystals.

Vitamin B Concentrate Japanese Patent 101,137

Rice bran or other similar vegetable material is extracted with methanol at 60° C. The solvent is distilled off in vacuo. The extractive residue contains a good percentage of vitamin B.

Detecting Cold Storage Eggs

By dipping eggs in lamp black, one can tell immediately whether they are

freshly laid or cold storage.

The test depends upon the fact that storage eggs are treated with an oil to preserve them. If it is a cold storage egg, the lamp black will cling readily to the outer shell, while the amount of lamp black adhering to a fresh egg is said to be negligible.

by exposure to air and to high temperatures than is the vitamin A in animal fats

tene is less readily destroyed

uble" (that is, dissolved in fats and not in water), it is not lost in cooking water, as are some of the "water sol-

uble", vitamins

Since vitamin A is "fat sol

VITAMIN DATA

Vitamins

Functions in the body

Good sources

Effects of various factors on the vitamin Long exposure to air, especially at high temperatures, may result in destruction of vitamin A, but it is not readily

Vitamin

ಹ barrier against the in-Maintenance of healthy membranes which provide Good health at all ages Successful reproduction It is essential for: Growth

vasion of bacteria

Its absence causes:

The surface covering in various parts of the body to may result in infection in the eye, in the respiratory low bacteria to enter, and break down. This may altract, and elsewhere

Cod-liver oil, halibut-liver oil, salmon and other fish oils Butter

Whole milk, cream, and cheese Carrots, pinnento peppers, spinmade from whole milk Liver and kidney Egg yolk

ach and other green leaves,

Usually, foods having a yellow or green color. Thus green leaves, yellow corn, and sweet potatoes are better sources than are blanched leaves, white corn, and white potaand tomatoes see

and green vegetables and fruits, may be changed to vitamin A in the body. Caro-

ing or canning processes
The yellow coloring matter,
carotin, which is found in
carrots and in other yellow

destroyed by ordinary cook-

Dried peas and beans Whole grains Nuts

Egg yolk Tomatoes Liver Milk Proper functioning of the di-Successful reproduction and

Green leafy vegetables

Good health at all ages

Normal appetite gestive tract lactation

(Anti-neuritic vitamin)

It is essential for:

Vitamin 2

Growth

Yeast

The beriberi, or polyneuritis

Its absence causes:

Ordinary cooking and canning processes do not destroy vitamin B readily, but since vitamin B is "water-soluble," much of it may be discarded if the cooking water or vege-table juice is thrown away

The addition of soda in cooking vegetables increases the Drying apparently does not destroy vitamin B destruction of vitamin B

Vitamin

(Anti-scorbutic vitamin)

It is essential for: Growth

Good teeth and healthy gums The maintenance of blood Good health at all ages

Fleeting pains in the joints, Insufficient amount may cause: sometimes mistaken for vessel walls rheumatism

Its absence causes:

Scurvy

Citrus fruits, raw or canned Tomatoes, raw or canned Raw cabbage Raw peppers

While they contain only fair amounts of vitamin C, raw and cooked potatoes may be important sources because apples, onions, and turnips, they are cheap and plentiful. Spinach

regulating the use of cal-cium and phosphorm

Good health at all ages Good bones and teeth

(Anti-rachitic vitamin)

It is essential for:

Vitamin

Growth

Rickets, which in turn may

Its absence causes:

the body)

cause permanent deformi-

ties of the bones

are not naturally good sources (as milk and bread) Vitamin D is now being intro-duced into some foods which by irradiation of the food or of some ingredient

Since vitamin D is not so sources need emphasizing widely distributed as other known vitamins,

Cod-liver oil, halibut-liver oil, salmon and other fish oils

Ultraviolet rays acting on the skin, either from sunlight or from special lamps (that is, carbon arc, quartz mercury-vapor lamp) Egg yolk

destroyed of the known vita-mins. Exposure to air, long the destruction of vitamin C cooking, and the addition of soda in cooking tend toward Vitamin C is the most readily

canning process tends to reduce the vitamin-C content of siderably, except in the case of the acid foods such as by the open-kettle Drying and storing foods tend to destroy vitamin C. The fruits and vegetables conmercially, air is excluded, and this process reduces the method, may lose more vita-min C than do commercially citrus fruits and tomatoes. When foods are canned com-Foods canned at home, espedestruction of vitamin canned foods cially

Ordinary processes of cooking do not easily destroy vita-Vitamin D may be somewhat more slowly destroyed by exposture to air than is vitamin A

Effects of various footows	the vitamin Ordinary cooking temperatures and exposure to air have little effect on vitamin G Use of soda in cooking has a destructive action on vita- min G	
	and	
VITAMIN DATA—Continued body Good sources	Fresh lean meat Liver and kidney Milk, fresh, evaporated, dried Buttermilk Salmon, fresh and canned Eggs Green leaves Tomatoes Yeast Wheat germ	THING STATE
VITAMIN Functions in the body	It is essential for: Growth Good health at all ages Prevention of symptoms similar to those of pellagra, such as digestive disturbances and skin lesions Its absence: Appears to be at least one factor in causing pellagra	
Vitamins	Vitamin G	

Egg Preservative British Patent 409,623

Eggs are coated with following:

Soft Yellow Paraffin
75 oz.
Tallow
5 oz.
Boric Acid
20 oz.

Destroying Yeast Spores in Soda Water Bottles

Soak for five minutes in 1% caustic soda solution at 45° C. and for 10 minutes in 2% caustic soda solution at 40° C.

Meat Curing Salt U. S. Patent 1,976,831

Mix together in an aluminum vessel Sodium Nitrite 1½ lb. Sodium Nitrate 1 lb.

Melt while stirring. Pour on metal plate to solidify. Pack in air-tight tins. For treating 100 lb. of beef use 1/4 oz. of above ground into 3 lb. of salt.

English Mustard, Prepared British Patent 412,967

Mustard flour is mixed with cold milk and water with 2% gum arabic and after ½ hour is sterilized by treating at 65-70° C. for 15 minutes, then cooled to 30° C.

Smoked Fish

It is hardly possible to furnish directions for smoking all species of fish, under all the varying weather conditions that will be encountered with the changing seasons. Only the general methods can be given here, as used on a typical variety under average conditions. This is intended as a guide, not an infallible recipe. To smoke fish successfully, experiment and use intelligence—altering the method according to the preference of markets (amount of salt and smoke flavor), the variety of fish, and weather conditions.

There are two general methods of smoking fish—hot smoking or ''barbecuing,'' and cold smoking.

Any fish may be ''hot-smoked'' or

"barbecued" but the following varieties are some of those to be preferred:

Butterfish Sailfish
Kingfish Spanish mackerel
Mullet Shad

Grouper Sturgeon is always hot smoked. Because of the keeping qualities of cold-smoked fish, certain varieties offer market possibilities for quantity production, such as:

Alewife or river herring

Shad Drum

Mullet

Red snapper

Redfish Grouper

Kingfish

Robalo or Snook

Squeteague (spotted trout)

Spots

In the first method the fish are laid three or four feet above a fire. and cured at temperatures from 150 to 200° F. The fish are wholly or partially cooked by this method, and therefore, no matter how carefully prepared, or how long smoked, will "keep" for periods of from a few days to a couple of weeks. If fish is to be preserved for any period of time, the cold smoking method should be used. In this process the fish are cured over a low smouldering fire at a temperature of 90° F., or less. The efficiency of the process depends on the drying action of the fire, which must be carried on at a temperature that will not cook the flesh. Fish may be given a short cold smoke, if preservation is intended for a few days only, or cured for several days if it is wished to "keep" them for some time. product is comparable to ham or bacon and should be cooked before using. The same general principles governing smoking, handling, and storing of cured meats should be followed in smoking fish.

A smokehouse for curing small lots of fish may readily be made, following instructions given here. Obtain a box or make one, about 6x3x3 ft. One end, that resting on the ground, should be removed. About 12 in. above this end a false bottom with auger holes at 2-in. intervals is built. This end of the box is set over a pit 2 ft. wide by 18 in.

deep.

A trench about 1 ft. wide by 1 ft. deep is dug from this pit for a distance of about 10 ft. The fire pit, a hole 3 ft. wide by 3 ft. long, by 18 in. deep, is dug at the end of this trench, which is then covered by sheets of galvanized iron, forming a chimney for the smoke from the fire pit to the smokehouse. If it is desired to build a more permanent house, terra cotta drain or sewer pipe may be used to connect the fire pot with the smokehouse. Cleats are nailed inside

the box on the sides, the first set about 12-14 in. below the top. The trays for holding the fish, or the ends of the smoke sticks rest on these cleats. A few holes should be bored for ventilation in or near the top of the house.

If mullet or Spanish mackerel are to be smoked, the following process is

recommended:

The fish should be split along the back just above the backbone, almost to the tail so that it will lay flat in one piece, leaving the belly portion solid. Clear out all traces of intestines, black skin and blood, taking special care to remove the coagulated blood and kidney just under the backbone. The head may or may not be removed, depending on the individual. If the head is cut off, the hard bony plate just below the gills should be allowed to remain, as it will be needed to carry the weight when the fish are hung on rods. If it is cut off the fish often pull loose and drop from the sticks.

After splitting and cleaning, the fish should be dropped in a brine made by adding two cups of salt to 4 gal. of water. They are left in this brine 30 minutes to soak out blood diffused through the flesh. At the end of this time they should be taken out, rinsed, and freed from any remaining traces of blood or other offal. Drain for a few minutes then drop each fish singly in a shallow box of fine salt, "dredging" it about, then picking it up with as much salt as will cling to it, and packing the fish in even layers in a tub or box.

The fish should be left in salt from 1 to 3 hours, depending on weather, size of fish, fatness, and length of time for which preservation is desired. The exact length of time must be determined by the smoker. When the fish are taken out of salt they should be rinsed in brine, scrubbing off all visible particles of salt or dirt. The fish should then be laid on chicken wire drying racks kept out of the direct rays of the sun, but located where a good breeze can reach Wire drying racks are desirable as the fish can dry on both sides. One side will remain wet, if laid on boards. The fish should be given about 3 hours drying, until a thin film is formed on the surface, before putting the fish in the smokehouse. If put in immediately after taking out of salt, the fish will be too moist, will require longer smoking, will not color and dry as well and will not have as good a surface.

The fish may be placed in the smokehouse on wire mesh trays, or hung on sticks or iron rods. In no case should any two fish touch as this will prevent the drying and penetrative action of the smoke. If hung on rods, more fish may be smoked at one time, and they will smoke better, with a clearer color. Trays, of course, give less trouble. Rods are run through the fish just under the hard bony plate at the neck, one rod on each side. Thus, each fish hangs from two rods. Twelve or fourteen fish may be hung on a set of two rods 3 ft. long.

The fire should be started an hour or two before the fish are put in the house. It should be low and smouldering. most any hardwood or wood other than pine may be used for fuel. Pine or other pitchy woods will give the fish a bitter taste. Some of the woods that may be used in the Southern States, are scrub oak, live oak, hickory, sweet bay, river mangrove, palmetto roots, button wood, and coconut husks. In smoking any one kind of fish, such as mullet, variety of flavor may be obtained through the choice of wood used in smoking. In addition to the woods listed above, orange wood gives a particularly pleasing flavor. Cypress may also be used. The fire should not give off too much smoke during the first 8-12 hours. A dense cloud of smoke should be built up for the balance of the process. The fire must be small and Two short chunks of woodsteady. about 2 ft. in length and the thickness of a man's arm are usually sufficient. The fire pit is kept covered with a sheet of metal to drive as much smoke as possible up into the smokehouse, and to keep the fire from burning rapidly. The fire must not be allowed to blaze up. The air should not feel warm on the hand if it is put in the smokehouse. The fish should be smoked for 24 hours, if they are to be kept for a couple of weeks, and for 4 or 5 days if it is wished to keep them for some time. The fire should not be allowed to die out at night or to be built up too large the last thing at night to make it last until morning.

After taking the fish out of the smokehouse dry for an hour or two in the air, then wrap in sheets of waxed paper, sprinkling a little fine table salt on each one, and store in tin or wooden boxes. Keep in a cool, dry place. If signs of mold appear, sponge off with vinegar and give the fish a short smoking

for from 3 to 6 hours.

Hot Smoking-German Method

The following method is recommended if it is desired to prepare a hot smoked

fish that can be used immediately without cooking. It will keep without molding or souring longer than other hot smoked fish.

Split, clean, and soak the fish to remove blood, as instructed previously. Then prepare a brine as follows: 2 lb. salt, 1 lb. sugar, ½ oz. saltpeter, 1 oz. crushed whole black peppers, 1 oz. crushed cardamom seeds. Make this up into a 90° brine, that is, one that will float a potato with a 10 d. nail stuck in it. Increase the amount of ingredients according to the quantity of brine you wish to make. The number of spices used can be increased in variety and amount. Various spice mixtures are used.

Put the fish in this brine for a period varying from 2 to 4 hours, depending on the size and thickness of the fish, amount of fat, and the taste of the individual. Some require a less salty taste than others. The exact length of time must be determined by experiment. Rinse off the fish in fresh water, and place on drying racks outside in a cool, shady, breezy place to dry for about 3 hours before putting in the smokehouse.

For the first 8 hours that the fish are in the house, give them a cool smoke in a dense cloud of smoke. Then increase the fire until the temperature is between 130 and 150° F. for 2 or 3 hours, or until the fish have a glossy brown surface. This partially cooks or "hot smokes" the fish. Wipe any moisture off the fish, and cool for a couple of hours before storing. Wrap in waxed paper and store in a cool dry place. Do not allow them to come into contact with ice, or store in wet cold.

In some cases the fish are brushed over lightly with vegetable oil (usually cottonseed) either just after finishing the cold smoking part of the process, or on taking out to cool. Another method of handling this fish after smoking is to cut the flesh up into fingers the length of a No. 2 can or pint glass jar. Skin and pack into the can or jar. Then add vegetable oil (cottonseed or olive oil, if you have it) until the spaces between the pieces of fish are filled and there is a layer of oil up to within an eighth of an inch of the top. Seal the cans or jars and store in a cool place such as an ice box until used. Under such conditions it should keep almost indefinitely. As this product is not "sterilized" the cans or jars should be thoroughly scalded before use. In some cases the oil is filled in hot and the containers sealed immediately.

Smoking Fish

Lake Herring and Whitefish

The process of smoking lake herring and whitefish is identical. If the fish are frozen when received at the smokehouse. they are thawed in the open air or better, by immersing and stirring them in a barrel of water of medium temperature. After thawing they are split down the belly to the vent, eviscerated, washed thoroughly, and pickled in butts or barrels, about 4 lb. of fine salt to 100 lb. of fish being scattered among them and sufficient brine of 90° salinity to cover them. Either dry salt or brine alone may be used, the former being preferred in warm weather and the latter during the winter. In case brine alone is used, some dry salt should be placed on top to strengthen the weak pickle floating at the surface. After remaining in the pickle for 10 to 16 hours, according to the strength of the pickle and the flavor desired, the fish are removed and strung on the smoke rods, 10 to 20 fish to each rod, according to its length and the size of the fish.

In stringing, some curers pass the rod through the body immediately below the nape bone, effectively preventing the fish from falling down in smoking, but also marring its apearance somewhat. more usual way is to pass the stick in at the right gill-opening and out at the mouth. Others pass the rod through the head near or through the eyes, and a few pass it immediately back of the throat cartilage. The latter leaves a neat appearance, yet it permits more fish to fall in the smoking process than when the rod is passed through the head or the shoulders. In some houses the smokestick is not passed through the fish, but instead a stiff iron wire, curved in "S" shape, is used to attach the fish to the stick, one end of the wire passing through the fish at the head or beneath the nape bone and the other hung over the smoke-stick. At Grand Haven, and to some extent in Chicago, Milwaukee, and one or two other places, the fish are secured by having stout smoke-sticks, about 11/2 in. thick and 21/2 in. wide; in the top of each, and about 1/4 in. from the edge, is driven a row of tacks or small wire nails at intervals of about 3 in., projecting about ½ in. above the surface. Ordinary cotton wrapping cord is tied to the wire nail at the end of each stick, and by means of this cord passing around each nail a single herring is held in place between each two nails throughout the length of the stick, the fish being placed with the back of the neck against the stick and the cord passing from one nail around the throat of the fish, entering under the gills on each side, and then around the next nail, and so on to the end. By having the stick of sufficient width, a row of small nails may be placed on each edge, so as to attach a row of fish at each side. This removes nearly all risk of the fish falling, and their appearance is not marred by holes through which the smoke-stick has been passed.

Some markets prefer the herring well smoked on the inside and to accomplish this the sides of the abdominal cavity are stretched open by means of small wooden sticks or tooth picks, either one or two sticks to each fish. This permits the smoke to permeate the stomach cavity better and results in a more durable article. In general, the western trade prefers the stomach cavity stretched open, while the eastern markets prefer them without the sticks; but there are exceptions. The smoked lake herring sold in Washington are mostly extended by means of a small stick, or, in case of large fish, by two small sticks.

The fish attached to the sticks are dipped in fresh water to remove surplus or undissolved salt, loose scales, etc., unless they have been rinsed before stringing, drained, and suspended in the smokehouse 4 to 8 ft. above the floor, and subjected to a gentle smoke for 4 or 5 hours. The door or damper is then closed, the fires spread or built up and the fish cooked for 1 or 2 hours according to the amount of fire, the height of the fish, and the particular cure desired. After cooling, which is accomplished either by opening the doors of the smokehouse or by removing the fish to the outside, they are ready for the trade. One hundred pounds of round fish, or 85 lb. dressed, yield about 65 lb. smoked. Ordinarily these fish keep one or two weeks, and even longer.

Lake Trout and Carp

Smoked lake trout and carp are prepared to a small extent in the manner already described for lake herring or whitefish.

Smoked Fish

Alewives, or River Herring

River herring or alewives are smoked in a number of localities, but principally in Maryland and Virginia.

In preparing these fish in the Chesapeake region they are washed in vats and scaled with a knife as soon as prac-

ticable after removal from the water. They are next immersed over night in strong brine, containing 12 to 14 lb. of Liverpool salt to each 100 lb. of fish, with some dry salt on top to strengthen the weak pickle that rises to the surface. The following morning the round fish are strung on smokesticks, the stick being usually entered at the left gill-opening of each fish and out at the mouth, as in case of hard herring or bloaters on the New England coast. The strings of fish attached to the stick are then dipped in fresh water to rinse them off, and after draining and drying for a few hours are suspended in the smokehouse about 6 or 8 feet above the fire, and exposed to a dense but cool smoke made of pine shavings or similar material for about 2 or 3 days. Care must be taken to prevent the fire from becoming too hot, thus causing the fish to crack at the lower end or possibly to fall from the sticks to the floor. Prepared in this manner the river herring will usually keep in good condition in the Chesapeake region for 30 days during the spring and for a somewhat less period in the summer. As the fish are not eviscerated before smoking the decrease in weight is small, 100 lb. of round fish yielding about 85 lb. smoked. The wholesale price is about 20 to 22 cents per dozen, according to the size and condition.

In Washington, Baltimore, and one or two other places the river herring are prepared in the following manner:

The fresh herring are scaled with a knife, gibbed like the pickled herring of Scotland, washed, and pickled for 3 hours in brine, about 20 lb. of Liverpool salt being used for each 100 lb. of fish. On removal from the pickle they are strung on small iron rods, the rod passing through the eye sockets of the fish, drained for an hour or so, and hung in the hogshead smokehouses, in the bottom of which a fire has been made of equal quantities of oak and hickory wood. The fish are dried for a few minutes and then the tops of the hogsheads are covered with old sacks or other suitable material. From time to time the fire is sprinkled with water to produce a vapor and the fish thus exposed to heat, smoke, and steam for about 3 hours, when they are removed and cooled and are then in condition to be eaten. Only oak and hickory should be used as fuel, as other materials do not produce the proper flavor. If the fire becomes too warm it should be smothered with oak or hickory

The process of smoking alewives com-

monly employed in the New England States differs from the Chesapeake process in a few minor particulars. The smokers are usually not so careful about removing the scales with a knife, depending generally on the frequent handling of the fish to scale them if cured soon after removal from the water. It is also customary in salting the fish to permit them to make their own pickle, the fish remaining in the pickle for 3 to 5 days. On removal they are soaked in fresh water for 5 to 6 hours and strung on hardwood sticks, the stick entering through the left gill-opening and out at the mouth. They are next rinsed, drained and dried for a short while and suspended in the smokehouse, where they are exposed to a smoldering fire of hardwood and sawdust for 3 to 4 days, when after cooling, they are ready for sale.

Shad

In the Chesapeake region and at various points along the coast small quantities of shad are smoked, usually in precisely the same manner as already described for river herring, or alewives.

Catfish

Being intended as a substitute, the catfish are smoked in identically the same manner as are sturgeon. The fish as received at the smokehouse are usually beheaded and eviscerated. They are skinned and cut into small pieces, weighing about 1 or 11/2 lb. each, and are pickled for 6 or 8 hours in tight barrels. This may be accomplished by rubbing the pieces with salt and placing them in the barrel either with dry salt scattered among them, or simply by placing them in the barrel with dry salt or with strong brine. On removal from the brine the pieces are rinsed by dipping in fresh water, to remove slime, surplus salt, etc.; they are then attached to the smokesticks and drained for an hour or so, and placed in the smokehouse where they are smoked for 7 or 8 hours in the same manner as sturgeon are treated. hundred pounds of dressed catfish yield from 65 to 70 lb. smoked, and the product sells usually at about 15 or 16 cents per pound. The total annual product of smoked catfish in the United States probably does not exceed 50,000 lb., and its sale is confined principally to those who are willing to accept a substitute because of its being cheaper.

At several points in the Mississippi Valley the small catfish are smoked whole, like lake herring. They are split to the vent and eviscerated, the head and in some instances the skin being left on, struck with salt in tight barrels, and smoked for a few hours in the manner described for lake herring.

Eels

Generally the eels are received at the smokehouse fresh, directly from the fisheries, but some are also received frozen from cold storage. In the latter case they are thawed by immersing them in water a few hours or by exposure to the open air. Some smokers "slime" the eels with salt; that is, rub the skin with a small quantity of fine salt to remove the slime therefrom. In dressing, the fish are split from the head to the vent and the viscera removed. It is desirable to continue the splitting down to the end of the tail sufficiently deep to remove the large vein along the backbone, but sometimes this may be pulled out without splitting the fish more than an inch or two beyond the vent. Few smokers, however, give attention to this item. The eels are immersed in strong brine from 13/4 to 71/2 hours, according to strength of brine, size of fish, and the desired flavor. This brine should be quite strong, about 20 lb. of Liverpool or other good salt being required for each 100 lb. of fish.

In New York the eels are usually pickled for 2 hours, while on the Great Lakes the length of the time is generally about 7 hours. On removal of the fish they are washed, bristle brushes being used by some smokers, while others simply dip the fish in water for removing the slime and surplus salt. A few smokers throw them in a tub of water and beat them with a net for several minutes to accomplish the same purpose. The eels are next strung on iron or steel rods one-third inch in diameter, the rod passing through the head of each eel, or through the throat cartilage and out the mouth, and hung in the open air for a few hours for drying. But if the atmosphere be moist or the saving of time necessary they may at once be placed in the smokehouse.

In New York, where small brick ovens are used, the fish are subjected to a mild smoke for about 4 or 5 hours until they have acquired the proper color, when the fires are gradually increased and they are hot-smoked or cooked for 30 or 40 minutes. At Buffalo and some of the other Great Lakes ports, the smoking is usually at an even temperature throughout and continues for 6 or 8 hours. Mahogany or cedar sawdust is used in New York for making the smoke, while hickory or white-oak wood is used for

cooking, the latter being preferred. Washington the eels are suspended in the hogshead smokehouses over a fire made of oak and hickory wood and dried for 20 minutes, when the hogshead is covered with sacking and thus hot-smoked for 3 or 4 hours, the fires being sprinkled with water from time to time to produce a hot vapor. The smoking must be carefully attended, for if the heat becomes too great the fish will curl up out of shape. A good test to determine whether the cooking is sufficient is the ease with which the skin may be separated or peeled from the flesh when the eel has been split.

The decrease in weight by dressing and smoking is about 35%, 100 lb. of eels yielding 65 to 75 lb. smoked. When eels have been pickled 6 or 8 hours they ordinarily keep 10 or 12 days; but when the salting has been only 2 hours, as is usual at New York, they are liable to mold after 5 or 6 days. Smoked eels keep a shorter length of time than almost any other smoked fish.

Eels are sometimes skinned before being smoked, the process being the same as described above, except that less salting and smoking is required, and it is also very difficult to keep them from falling down off the rods in the smoke-house

Salting (Including Corning) River Herring

The fish are usually taken from the boats on the day they are caught, but in some cases not until the third or fourth day. All handling of the fish is with scoop nets. When taken from the boats, they are spread upon the wharf for cutting. Sitting on a low inclined seat with his knees on the wharf, the cutter removes the head and belly and scrapes out the roe and viscera, the cut fish being placed in a basket and the roe in a bucket. The fish are then dumped into the washing vats. These are 12 ft. long by 6 ft. wide by 3 ft. deep of 2 in. pine. In some the bottom is inclined about 30° to one side, with a horizontal false bottom of slats above the incline. Scales, dirt and other washings settle down in the deep angle of the bottom and are drawn off with the wash water through two flood gates without loss of time. Others still employ flat bottomed vats with resultant loss of time in clean-

ing.
The fish are agitated in the vats (which are kept filled with water) for about 10 minutes to thoroughly wash them and then scooped out with dip nets

into slat cars holding about 1200 fish, in which the fish drain as they are transported to the salting vats. The latter are 10 ft. long by 6 ft. wide and 24 to 30 in. deep built of 2 in. Virginia pine. The salting vats contain saturated brine to a depth of 4 in. As each car of fish is dumped into the brine, additional salt is added, the amount depending upon conditions of temperature of fish, etc., with which the skilled packer is fully conversant. When full, the vats contain from 12,000 to 15,000 fish (about 4000 lb.). The fish should be roused once each day while striking. After each rousing, the fish are tamped down lightly and top dressed with a thin layer of salt.

Corning

Early in the season most of the packers in the lower Potomac corn their herring for immediate consumption. This method is usually followed for about 6 to 10 days from April 1. The earliest caught fish are kept in the brine from 12 to 48 hours according to temperature. Fish brined 12 hours when the temperature is from 40 to 50° F. should keep for ten days. After brining, the fish are taken from the vats and spread on the floor, covered with salt and the salt and fish thoroughly mixed, after which they are packed in sugar barrels and immediately shipped to the trade. No fish are corned after the temperature rises above 60° F.

Hard Cure or Tight Pack

Herring intended for storage are kept* in the brine for 7 to 10 days according At temperatures from to temperature. 50° to 60° F. 9 to 10 days is sufficient; if from 60° to 70° F., 7 to 9 days will cure them satisfactorily. After the fish are cured, they are taken from the brine and piled on the draining floor to a depth of from 1 to 4 ft. according to available space and allowed to remain there from 4 to 10 days according to the demand for the space. The fish are then weighed or counted (weighing is most accurate) and packed in the barrels, the first layer backs down, the balance backs up with from 2 to 21/2 lb. of salt to the layer. A properly packed barrel should contain 160 lb. of fish and 40 lb.

Salted Fish

Considerable trouble has been experienced in salting fish in warm climates. The methods followed commercially in other regions have not produced a

product of good quality, and the directions given generally for salting small quantities, or for the home curing of fish have not always proven satisfactory.

If attempts are made to preserve fish by "pickling" or curing in brine, in a warm climate, the product will either turn "rusty" and sour, spoiling in a short time, or if the quality is good at first the fish soon deteriorates. The best method for curing fish in this region is "dry-salting." That is a combination of salting and drying. If the fish are handled carefully, and directions given below followed closely, a high quality product that will not spoil nearly as rapidly as salted fish now prepared, can be produced. But if instructions are not followed, it is useless to expect much.

In the first place the fish must be absolutely fresh. Do not try to save fish that may be stale, by salting. The fish should be bled, when caught, to drain out all blood possible. Blood decomposes much more easily and quickly than flesh. Fish will keep longer if blood is not diffused through the flesh. should be thoroughly cleaned as soon as possible. Fish should not be handled roughly in taking out of the net or while in the boat. If fish are piled in heaps, walked on or forked roughly, they will be of inferior quality and spoil much more readily than they would otherwise. Fish should not be left under the direct rays of the sun in an open boat. A tarpaulin should be rigged above the

Mullet and Spanish mackerel are among the best fish for dry-salting, for many reasons, a few of which are: they are split more easily, the loss of weight is less in splitting and cleaning; they are two of the commonest southern fish, and obtained more easily and cheaply. Using this outline as a guide, however, many other varieties of fish, such as grouper, sheepshead, alewives or river herring, spot, croaker, and drum, may be cured successfully, with the resultant product of good quality.

Most fish should be split along the back, just above the backbone, taking care to leave no flesh on it. The fish are split "mackerel style." That is, they must lay flat in a single piece, leaving in the backbone. When the knife is drawn toward the tail it must not go clear through the skin, so that the fisk will be in two pieces near the tail. The head may or may not be removed. In splitting Spanish mackerel and other fat fish the backbone is cut out nearly to the tail, where it is broken off. In

cleaning, remove all traces of blood from under the backbone and clear away all the black skin. A wire brush should be used for the blood. "Black skin" is best wiped out by a piece of canvas or gunny sack. If the head is left on, clean out all traces of gills. All cleaning must be done thoroughly and carefully.

When the mullet or mackerel are cleaned they should be rinsed, then dropped in a tub of light salt brine (2 lb. of salt to 5 gal. of water), the fish should be left here to soak 30 minutes. The principal object of brining is to remove traces of blood from the cut flesh. It also "cuts" slime and is better for washing than water. Never use sea water from around a fish house, dock, or near shore. It is invariably contaminated and increases likelihood of spoilage.

Score with a knife under the backbone and then longitudinally through the flesh on the other side. After the fish have soaked 30 minutes take them out, making sure that each one is properly cleaned. Drain them for 15 minutes. If salted at once the excess moisture will

require more salt.

Use a "dairy fine" ground mined salt. Ordinary sea salt is more apt to cause reddening. Coarse salt is not as good as a fine salt. Pour the salt into a shallow box about 2 ft. square. Dredge each fish in this salt, rolling it about 2 or 3 times and rubbing salt into the slashes. Pick it up with as much salt as will stick to it. Scatter a thin layer of salt on the bottom of the tub or box used for salting. Then lay in the fish in an even layer, flesh side up. Be sure that no two pieces of fish touch without salt between. Scatter a little salt on top. Continue this until all the fish are in salt. Each layer should be laid in at right angles to the preceding layer. The top layer should be weighted down, to keep the fish under the surface of any brine The top layer should also be formed. packed skin side up. Use about 1 part of salt to 3 of fish.

The salting shed should be light, open, airy, and cool as possible. The mullet will have absorbed enough salt for curing purposes in about 36 hours. Mackerel should be in salt about 48 hours. At the end of this time take the fish out of the salt and scrub them in a brine of the same strength as used in cleaning to remove all excess salt and dirt. No traces of salt should be visible on the surface. After draining 15 to 20 minutes, the fish are ready for the drying racks. These are frames of wood, cov-

ered with chicken wire and standing on

legs 3 or 4 ft. high.

The drying racks must be placed on dry ground, preferably covered with gravel. Oxidation or rusting sets in immediately if drying is carried on under the direct rays of the sun. But if fish are kept shaded in a breezy location they will dry well with a clear color. For this reason drying is best done in the shade under a roof without walls, so located that as much of a current of air as possible will pass over the fish. The fish are laid out skin side down but are turned 3 or 4 times the first day.

The fish are gathered up and placed under shelter at night to prevent spoilage through dampness. If left spread out in the open at night, they will sour and mold. The time required for drying depends on weather conditions during the drying period, and on the size of the fish being cured. The exact time must be determined by the person curing the fish. For mullet it should average about 4 days; Spanish mackerel, 5 days. The more the fish are dried, the less danger there will be of reddening or rusting. When the surface looks dry and hard, and if the thumb can be pressed into the thick part of the flesh leaving no impression, the flesh can be considered as cured.

In weather where air-drying is impossible, or in climates too humid for this process, the following method may be used. When the fish are "struck through" or have absorbed enough salt for curing purposes, they should be taken out of salt, scrubbed off in brine, then piled in stacks, flesh side down. These stacks should be heavily weighted down in order to press moisture out of the fish. After 10 to 18 hours in the stack the fish should be repacked in dry salt with the top weighted down, and put in storage in a cool dry place.

Store the fish in wooden boxes lined with waxed paper. Scatter a little dry salt between each layer of fish—about 1 lb. of salt to 10 lb. of fish. Store in as cool and dry a place as possible. If signs of rust or mold appear, scrub the fish off in brine and dry in the air for

a day or two.

Reddening of salted fish is a form of bacterial spoilage caused by the salt used in curing. Contrary to popular belief, salt is not strictly an antiseptic, and certain types of bacteria live and thrive in a salt medium. Salt most apt to be contaminated is that obtained by evaporation of sea water. Several types of salt used extensively in fish curing are apt to be thus contaminated. In salting fish

every effort should be made to use a salt as pure and high in grade as possible. It is advisable to heat salt and bake it thoroughly before using. If, however, reddening appears at any time, all tables and other equipment used in salting should be thoroughly disinfected. Unless every effort is made to keep the salting equipment clean, the use of sterilized salt or other precautions will be useless as the fish can be contaminated through unclean equipment. After curing, the fish should be stored in the coolest place possible, as the salt reddening bacteria grows best at a warm temperature. At first signs of reddening the fish should be removed, washed thoroughly in pure salt brine, and given a few hours careful drying and repacked with a thin layer of dry salt between each layer of fish, using from 10 to 15 lb. of salt to 100 lb. of fish. Reddening is most apt to appear in fish stored in pickle (brine) and held in a warm place. It will remain in good condition longer if packed in dry salt and held in as cool a store room as possible.

Canning Alewives or River Herring; Roe and Buckroe

The following method of canning alewives has proved quite satisfactory. The fish are cut, washed, and placed in the salting vats in the same manner as if intended for salt curing. After 12 to 14 hours they are removed from the vats and washed in an abundance of lukewarm fresh water. During the washing they are trimmed, the balance of the fins and scales being removed. They are then cut to can size and placed in the cans, after which they are processed for 55 minutes at 244° F. for No. 1 cans and 60 minutes for No. 2 cans.

Herring roe intended for canning is collected in buckets as the fish are cut and washed in fresh water in special trays, blood and adhering particles of entrails being removed. The roe is then put in the cans. As it swells considerably in processing, the cans must not be entirely filled. If of the sanitary type, the cans are filled to within about threefourths of an inch of the top with roe and then filled to the edge with cold salt brine, about 1 lb. of salt to 8 or 10 gal. of water being used to make the brine. The brine is added solely for seasoning. The cans are immediately capped and placed in the processing baskets. If solder-top cans are used, the filled cans are placed in the exhaust box. removal from the exhaust, the necessary

air space is provided for by pressing the roe down with a plunger. Material clinging to the groove where the solder is to be applied is removed with a brush and the cans are capped and tipped. canned roe is processed in a closed kettle for 45 to 55 minutes at a temperature of 240°-245° F. The milt roe may be canned in the same manner as the roe except that the cans can be more completely filled, as this product does not swell in the processing. As the quantity of brine used in this case will be somewhat less, it should be made correspondingly stronger.

Note: In canning the fish, they should be drained of superfluous water before they are placed in the cans, and no water added to can contents. That the fish may retain their shape in the can and stand transportation, the cans should be well filled. The shrinkage of the fish in processing must be taken account of in filling the cans.

Canning Clams (Alaska)

The first operation is the removal of the clams from the shells. This is done by immersing them in boiling water, either in vats especially designed to receive the wire baskets in which the clams are placed or the clams are passed through the water on an endless belt. After remaining in the water several minutes they are thrown on a table and the shells fall away from the meat. The clams are then passed on to women workers, who open the stomachs and necks, remove the sand and sediment therefrom and sever the black part of the neck. The cleansing process is continued by placing the meat in a cylindrical perforated washing machine, which revolves automatically half a turn both ways in a tank filled with water. Any sediment that may have remained after the hand operations were completed is thus removed. The clams are now ready to be canned and are taken directly to the filling tables if whole clams are packed, or to the grinder if the minced variety is desired. The cans are filled by hand with both meat and juice, after which they pass through the topping and sealing machines and are sealed. The process is completed by cooking the canned product in retorts at a temperature of about 245° F. from 1 to 11/2 hours, depending upon the size of the container used. The juice which is thrown off in the process is used in preparing the finished product, the surplus being sealed in cans.

Anchovy Paste

Anchovy paste from sprats may be made as follows: Sufficient for a peck of sprats—2 lb. common salt, 3 oz. bay salt, 1 lb. saltpeter, 2 oz. prunella, and a few grains of cochineal, pounded well together in a mortar; into a stone jar place first a layer of fish, then of the pounded ingredients, and so on until the jar is filled; press them hard down and cover closely. After 6 months they will be ready for use.

Note: Persons using such preservatives as saltpeter should consult the Bureau of Chemistry, Washington, D. C., to determine whether they are using an amount in excess of that held to be proper under existing law.

Anchovy Butter

Take 1 part of anchovies which have been beaten to a paste, and pass through a sieve; add 2 parts of butter, and spice to suit. Cayenne pepper or paprika may be used to advantage.

Anchovy Essence

Anchovy essence can be made with either canned or bottled anchovies. Take the fish, and rub to a pulp in a mortar, and then pass through a fine sieve. To ¼ lb. of anchovies add ¼ lb. of water; boil for 15 minutes, and strain; then add ½ oz. of salt and ½ oz. of flour, and the pulped anchovies. The mixture is allowed to simmer over the fire for 3 or 4 minutes. After the preparation is cool add 2 oz. of strong vinegar. The product should be bottled in small bottles and tightly corked and covered with bottle wax.

Anchovy Paste

Prepared by taking 1 lb. of anchovies, 1 lb. of water, and $2\frac{1}{4}$ oz. of salt and $2\frac{1}{4}$ oz. of flour; add a small quantity of cayenne pepper (say $\frac{1}{10}$ oz.), a small quantity of grated lemon peel, and $\frac{1}{2}$ oz. of mushroom catsup.

Anchovy Sauce

Take 3 or 4 anchovies, and chop them fine; add 3 oz. of butter, 2 oz. of water, 1 oz. of vinegar and 1 oz. of flour. Melt the butter over a water bath, add the water and the vinegar, and lastly the flour and the anchovies; stir until the mixture is thick, then rub through a wire sieve. This preparation should be kept on ice, and will not keep indefinitely.

Mushroom Catsup

Upon a suitable quantity of the fresh mushrooms sprinkle salt (about 1 to 4 of the fungi), and after 3 days squeeze out the juice. To every gallon of juice add black pepper, ginger and cloves, of each ½ oz.; pimento, 2 oz.; mustard seed, 2 oz.; and a sufficient quantity of salt. Boil for 5 minutes and set aside to settle. Strain after 7 days.

Christiana Anchovies

In the preparation of Christiana anchovies many methods and flavoring ingredients are used, depending on the skill and ideas of the curer and the markets for which the preparation is intended. The following is one of the most popular processes:

The fresh sprat or anchovies are immersed in brine for 12 or 18 hours, 15 lb. of Liverpool salt being used for each 100 lb. of fish. On removal, the fish are drained in a sieve and then loosely packed in a barrel, with the following ingredients, which have previously been finely crushed and well mixed: 4 lb. of Luneburg salt, 6 units of pepper, 6 units of sugar, 6 units of English spices, 1 unit of cloves, 1 unit of nutmeg, and 1 unit of Spanish pepper. The anchovies remain saturated with these ingredients for 2 weeks, when they are repacked tightly in kegs or barrels, being carefully arranged in layers, with the backs downward. A quantity of the ingredients above mentioned is sprinkled over each layer, with the addition of a few cut bay leaves or cherry leaves. At the bottom and the top of the package is placed two whole bay leaves, but before the top leaves are laid on, brine is poured over the fish. The barrels or kegs are then coopered and rotated daily for the first few days, and after that every other day for 2 or 3 weeks.

The following process is also used to some extent.

The fish are salted for 24 hours and next immersed in sweetened water, 20 parts of water to 1 part of sugar being used. The fish are then packed with a mixture of Luneburg salt with 90 units or parts of allspice, 60 units of pulverized sugar, 19 units of whole peppers, 15 units of cloves, an equal quantity of nutmeg or mace and of hops (Origanum creticum), and some bay leaves.

The following is a choice method of preparing "Matjeshering" in Germany:

Fresh full herring, both spawners and milters, are well washed, and the gills,

stomach, and intestines are removed in such a way as not to necessitate cutting the throat or abdomen, this being accomplished by pulling them through the gill flap. The fish are next immersed for 12 or 18 hours in a 7% solution of whitewine vinegar, from which they must be removed before the skin becomes flabby and be wiped dry and covered with a preparation composed of 2 lb. of salt, 1 lb. of powdered sugar, this quantity being sufficient for 75 herring. The fish are then packed in a barrel which is sealed. When there is not sufficient brine to fill the barrel, additional should be made of 1 part of the above mixture and 4 parts of water which has been boiled.

Spiced herring (Gewurzhering) are prepared in Germany in the manner above described, with the addition of spices mixed with the salt. The spices commonly used consist of 1 part of Spanish pepper, 5 parts of white pepper, 4 parts of cloves, 2½ parts of ginger, an equal quantity of mustard, and a particle of mace and of Spanish marjoram, with a few bay leaves scattered between the layers.

Smoked Pork Sausage

Formula.—Meats: 100 lb. strictly fresh pork trimmings, 85% lean and 15% fat

Seasoning:

Salt	21/2 lb.
Ground White Pepper	10 oz.
Granulated Sugar	4 oz.
Ground Nutmeg	1 oz.
Ground Ginger	$\frac{1}{2}$ oz.
Nitrate of Soda	2 oz.

Nutmeg and ginger may be omitted and sage substituted. Some classes of trade prefer this product with only salt, pepper, sugar and nitrate of soda in the

seasoning formula.

Processing.—Inspect pork trimmings to see that they are fresh and lean. It may be necessary to re-trim, removing blood clots, gristle and hair. Proportion of fat and lean should be closely watched since fat has a tendency to render out in the smokehouse and soften the product. Grind pork through 5/32 or 4-in. plate of the hasher, first making sure knives and plates are sharp. Some packers use a rocker entirely for pork sausage.

Place meat in mixer and add seasonings. Mix seasonings and meat for about 5 minutes or until ingredients are thoroughly intermingled. At the time seasoning is added a small quantity of

crushed ice (not more than 7 or 8 lb. per 100 lb. of meat) may be used.

Stuffing.—After seasonings, meat and ice are thoroughly mixed, the product goes to the stuffing bench where it is stuffed in medium hog casings. Link in double links, 3½ in. in length, knotting ends of casing to prevent meat dropping on truck or floor. Trim off all scrap ends of casings on the outside of knot, but be sure scraps do not get mixed in with the meat.

Carefully puncture casings to prevent air pockets between casings and meat. Sausage must be hung on a truck as fast as it is linked. When truck is filled, put it under an overhead cold water spray for several minutes to thoroughly remove grease and sediment from outside

of casings.

Scrap meat on the bench should be handled promptly and mixed with meat stock in the truck. It should not remain on bench for any length of time as it

deteriorates rapidly.

Cooling.—After stuffed sausage has been sprayed it is taken to cooler and spread on trucks or in hanging sections and allowed to hang overnight at a temperature of 36 to 40° F. Product is removed from cooler the next morning and allowed to remain in natural temperatures for about 2 hours.

Smoking.—Then it is placed in the smokehouse at a temperature of 115 to 120° F. and carried at this temperature for about 3 or 4 hours. It does not re-

quire a heavy smoked color.

After smoking it is placed in the cooler at a temperature of 45 to 50° and allowed to hang for 2 to 3 hours until thoroughly cooled. Then it is packed in cartons if it is to be shipped promptly. This product should be manufactured only as needed.

Pork Sausage

Meats: Cali Butts 45 lb. Selected Ham Fat 55 lb. Seasoning: Salt 1¾ lb. Fine White Pepper oz. Fine Sage 2¾ oz. Cardamom 1/3 oz. Savory OZ. Marjoram OZ. Ginger OZ. Sugar

Put ham fat on rocker with 3% ice for 8 minutes, then add seasoning and lean meat and rock for 10 minutes more, making 18 minutes altogether. Meats are all fresh and in small pieces. When rocking is finished fat must have the appearance of half the size of a coffee bean.

Another meat formula for breakfast

sausage is as follows:

Shoulder Fat Pork
Trimmings 25 lb.
Pork Butts Trimmed 25 lb.
Lean Pork Trimmings, 40%
Lean (No Belly Trimmings) 50 lb.

"Skinless" Pork Sausage

Sausage meat for this product is stuffed in "NoJax" or similar casings, linked usually in about 4½-in. lengths, and handled and peeled in same manner as skinless frankfurts.

Following are two formulas for "skin-

less'' smoked sausage:

For formula No. 1 use, per 100-lb. batch:

Lean Pork Trimmings, Cured 60 lb.
Regular Pork Trimmings,
Cured 20 lb.
Lean Beef, Cured 20 lb.

Pork is ground through 1/4-in. plate. Chop beef very slightly so it will act as a binder and then add to pork in mixer. Care should be taken that no excess moisture is added as it will produce sourness in finished product. Mix well and season with proper amounts of salt, pepper and whatever other seasonings are desired.

Ready prepared seasonings or specially prepared seasonings as manufactured by reputable firms will assure convenience and uniformity in making this

product.

Stuff mixture in 1½-in. "NoJax" or similar easing. Smoke in a cool house for 3 hours at 130° F. Then cook at 160° F, for about 10 minutes. Cooking is usually done in a steam house to prevent smearing. Sausage should be placed before a fan following cooking to dry off casing. This aids in prevention of any mould or bacterial growth.

Formula No. 2 uses, per 100-lb. batch:
Cured Pork Cheeks 50 lb.
Cured Regular Pork Trimmings 50 lb.

This formula is prepared in same manner as No. 1. Product must not be chopped too fine or cooked too much to prevent pork from becoming smeary and spoiling its appearance. Sausage should not be peeled or packed in boxes until ready for shipment.

Italian "Hot" Sausage

A good formula for this product is as follows:

Beef, Free of Sinews 60 lb.
Pork Trimmings (Half Regular and Half Lean) 40 lb.

Chop meats through the 1-in. plate and mix with following:

No. 3 Can Pimientos, Juice and All, Chopped to a Paste 1

Straight Ground Chili

Pepper 1½ lb. High Grade Paprika 1 lb.

If fresh meat is used in making the product 2 lb, of salt should be added. If meat is cured, the additional salt is not necessary. Also add:

Ground Caraway 1 oz.
Coriander 2 oz.
Celery 1 oz.
Nutmeg 2 oz.

After a thorough mixing, run the product through 32, 16 or 1-in. plate, depending upon fineness or coarseness of

meat desired.

Stuff mixture in hog or manufactured casings, linked 6 to pound. This allows serving two sausages on average plate lunch. Put sausage into cook tank with water at 160° F. and let temperature drop back to 150°. Cook for 30 minutes or until an inside temperature of at least 137° is obtained.

This sausage can be smoked right after it is stuffed, smoking for half an hour

in a cold smoke.

Any good bologna or frankfurt meat formula can be used for this sausage, cutting the meat coarser if desired and seasoning highly, with seasonings such as those suggested in the above formula.

Another meat formula which might be

used is as follows:

Beef Chucks 70 lb.
Pork Cheek Meat 20 lb.
Back Fat Trimmings or
Shoulder Fat 10 lb.

Grind beef and pork cheeks through the 1/4-in. plate; back fat trimmings through 3/6-in. plate.

Head Cheese

The following formula can be used to make an attractive product which is strictly a head cheese.

Meats:

S. P. Pork Tongues	60 lb.
S. P. Pork Snouts	20 lb.
Pickled Pork Ears	10 lb.
Pickled Pork Rinds	10 lb.

Seasoning:		
Ground White Pepper	4	oz.
Caraway Seed	2	oz.
Marjoram	1/2	oz.
Ground Cloves	1/2	oz.

Prepared seasonings may be used if desired, such as those made by reputable seasoning manufacturers, to facilitate convenience in handling and uniformity of product.

Cook each kind of meat separately in nets, at 212° F. as follows:

1½ hr.
2 hr.
1½ hr.
1¾ hr.

Grind skins through 1/8-in. plate of hasher. Snouts and ears should be put through 1-in. plate. These should be rinsed several times with warm water to remove surplus sediment and fat.

remove surplus sediment and fat.

Remove gullet bones from pork tongues after cooking. Cut each tongue crosswise 3 times, making 4 approximately equal pieces, so that tongues will pass through valve of stuffing machine.

Put all meats together in a box truck, adding seasoning, jelly water and salt to taste. Not much salt will be required, as all meats used are pickle-cured. Use the hot meat liquid in which meats were cooked, and mix thoroughly.

Stuff tight in hog stomachs or manufactured casings. Fasten carefully and cook 1½ hours at 170° F. Wash clean and put into cooler at about 36°, or keep in ice water, to chill thoroughly before packing. Product must be clean and free of grease before packing and sale.

Some sausage makers add pimentoes or green peppers to give eye and taste

appeal to their head cheese.

Curing and Smoking Frankfurters

Curing is best done by dry-curing hashed meats, by emulsion curing, or by a combination of both. In dry-curing hashed trimmings use per 100 lb. of meat, 3 to 3½ lb. of salt. Nitrate or saltpeter should never exceed 3 oz., while nitrite should never exceed ¼ oz. per 100 lb. of meat. A mixture of these is still better, namely ½ to ¼ oz. of nitrite and 2 to 2½ oz. of nitrate or saltpeter. The same proportions hold for the emulsion cure. Dry cured hashed trimmings may be used after 2 to 3 days, but they may also be kept 7 days. Emulsion cured meats are put through the fine cutter, and so cure rapidly. Thus they must be used promptly.

Every sausage maker knows that good

muscle meats make good sausage and that cheeks and other such meats do not make sausage of quite as high a class. Less ice should be used in the summer than in the winter. For winter about 60 lb. of ice can be used per 100-lb. block of meat, but only 40-48 lb. should be used in the summer for first grade frankfurters. Less ice can be used with second and third grade frankfurters.

Frankfurters should be properly cured before smoking. If the emulsion cure is used in whole or in part, the meat or the sausage should be held a while for the cure to develop. Part of this may be done in the smokehouse. The smoke should start cool (about 90° F.) and finish at 130-135° F. for frankfurters and 140-145° F. for Vienna style frankfurters. For other smoked sausage the finish may be at up to 175° F. Cooking should follow promptly and the two operations should really be considered as one. Vat water should be 160°-165° F. while in the spray cooking process the water may be 180° F. Cooking should proceed until the temperature at the center of the meat is at least 140° F. while 148° F. gives better color and many believe it gives better texture and flavor.

German Ham

Since these hams are not cooked before they are eaten, all packers operating under federal inspection must follow B.A.I. rules for uncooked pork in making them. The way they make them in Germany is as follows:

Only hams with a pink meat color are chosen. They should weigh about 18 lb., and are long cut with some of loin end on. Hip bone should be removed.

For curing use a mixture of 25 lb. of salt and 4 oz. of sodium nitrate, or prepared curing mixture. This mixture is rubbed into the ham, especially the skin side, for about 5 minutes. Press some of salt into leg bone at cut. Place hams in a vat, and on each layer add enough of curing mixture so that all parts are lightly covered with it.

When vat is full it should be covered with boards with a weight on top. Curing will take 28 days at not less than 38° F. Repack 3 times during this period, so that top layer goes on bottom. Rub hams over again at each repacking.

Rub hams over again at each repacking.
At end of 28 days take hams out of vat and lay on floor in same temperature for 14 days, sprinkling curing mixture very lightly between each layer. At end of this period wash hams in warm water and hang in dry-room for 2 to 3 days.

Then smoke in a very cold smokehouse for not less than 6 weeks. In Germany these hams are sometimes smoked for 6 months.

Careful handling in cure will yield a tender product. Packers preparing this type of ham for the first time should cure only a small batch. In this way they can watch smoking and curing closely.

Bologna

To make and cure bologna in the silent cutter one sausage expert advises the use of all fresh meats, as follows:

Beef Chucks	70 lb.
Pork Cheek Meat	20 lb.
Pork Back Fat Trimmings	
or Shoulder Fat	10 lb.

Grind beef and pork cheeks through the 1/2-in. plate; back fat trimmings through 3/2-in. plate. Put beef and pork cheeks in silent cutter and add cure, as follows:

Salt	3	lb.
Sodium Nitrate	2	oz.
Nitrite of Soda	1/4	oz.
Sugar	6	0Z.

and proceed as if using cured meats.

Add ice and water up to 20 lb. per 100 lb. of meat, and chop for 3 minutes.

Then add pork back fat and seasonings:

F	- 4
Ground White Pepper	6 oz.
Ground Allspice	1 oz.
Coriander	2 oz.
Ground Nutmeg	2 oz.

Chop 2 minutes more. Then put in a meat truck or pans not over 6 in. deep, and hold in cooler at 36 to 38° F. over night or about 12 hours. Next morning stuff and let sausage hang in room temperature for 1 to 2 hours. Then smoke, slowly at first, gradually increasing temperature from 120 to 145° F. Cook 45 minutes at 160° F.

This method has the advantage of saving a lot of labor, decreases inventory holding and produces a fine, tacky product.

Non-Discoloring Salami

Discoloration is usually due to curing methods. To make either hard or soft salami, meat should be cured as follows:

Use 234 oz. of sodium nitrate for each 100 lb. of meat. Beef requires 3 lb. of salt and pork 214 lb. for each 100 lb. of meat cured. Run meat through 1-in. plate with above curing materials and then cure for at least 8 days at a temperature of about 40° F. Then place in

mixer, add 9 oz. sugar and 6 oz. of pepper, and mix pork and beef together. Grind mixture through desired plate, either ¼ in. or % in.

Stuff material tightly in large hog bungs, beef middlings or manufactured casings, as tightly as casing will stand. Hang in a dry chill room for 4 days. Then remove to sausage kitchen and hang for at least 6 hours so it will be raised throughout to room temperature before it goes to smokehouse. It may either be smoked through or smoked 12 hours and finished in cooker.

"Smoked through" means about 24 hours at slow smoke at 90 to 100° F. Then gradually raise temperature to about 140° so that product will have a 137° temperature at center when finished. Remove from smokehouse and rinse off with cold water; allow it to cool before placing in chill room.

Meat from full grown animals should always be used for hard sausages, such as jumbo shoulder trimmings and large beef chucks with all sinews removed.

A good meat formula for salami is as follows:

Lean Pork Trimmings	50 lb.
Medium Lean Beef Chucks (Free of Sinews)	35 lb.
Back Fat	15 lb.
There meets should be sured	nacordina

These meats should be cured according to directions given previously.

The product may be seasoned with:

Crushed Garlic	11/4	oz.
Sugar	9	oz.
Brandy Flavoring	5	oz.
Ground Anise Seed	1	oz.
Ground Cardamom	1/2	oz.
Maple Flavor	3	tbsp.

Coloring and Flavoring for Meats British Patent 425,567

Hæmoglobin, Defibrinated	100	oz.
Sodium Nitrite	5	oz.
Sodium Nitrate	$1\frac{2}{3}$	oz.
Water	100	oz.

Stir well for a few hours. Spray dry or vacuum dry. 1% of this product is used on meats.

Preserving Color of Meat U. S. Patent 2,009,587

By coating freshly cut meat surfaces with a glycerin-gelatin-water solution containing a small amount of essential oil, the natural fresh color and appearance of the meat is maintained.

Various essential oils, such as oil of cloves, may be used, or a mixture of oil of black pepper, coriander and all-

spice.

One typical formula for such a solution that has been found satisfactory consists of 57% water, 25% glycerin, 18% gelatin, and substantially 0.1% of essential oil. This solution may be applied with a brush or spraying device on cloth placed over the cut surface of the meat.

The entire piece of meat may be wrapped in fabric such as export beef cloth or the fabric may be applied only on the cut surfaces. The coating is then allowed to congeal. The glycerin, being hygroscopic, preserves the gelatin in a flexible condition, thus avoiding cracking. The essential oil acts as a germicide, while the gelatin acts as a hermetic seal.

Export beef cloth has been found superior to other fabrics for keeping the preservative solution in contact with the

Preserving Vegetables and Fish Dutch Patent 34,553

A procedure for keeping fruit, vegetables, etc., in a fresh condition has been devised. It is especially adapted for the prevention of mold, fungi, and other micro-organisms developed during storage. The procedure consists in rendering the air of the storeroom slightly alkaline, so that moist indicator paper showing a color change at pH == 7.5 is affected on introduction into the chamber. In order to render the storeroom alkaline, materials which furnish volatile, alkaline substances are burnt slowly.

Preventing Mold on Stored Meats

The humidity of the cooler should be 90 to 92% and the temperature 38-39° F. Ozone is introduced until it is present in 2.3 to 2.7 parts per million. This is continued for 2 hours and again for 2 hours after a lapse of 12 hours. After an interval of 30 minutes, workmen can safely enter the room.

INKS AND MARKING COMPOUNDS

Ink for Documents	Pour this into:
Gallie Acid 5 g.	Water 180 g.
Borax 0.5 g.	Indigo Carmine, Paste in
Pierie Acid 2 g.	Water 36 g.
Ammonia 20 g.	Wood Vinegar, Crude 15 g.
Water 50 g.	Dye for Black Writing: per 1000 cc.
Dissolve with warming and stirring.	Ink add: Phenol Blue 3F 1.8 g.
Water 50 g.	Phenol Blue 3F 1.8 g. Ponceau RR 1.2 g.
Caustic Potash 1 g.	Aniline Green D 1.2 g.
Boil and stir the mixture until pale	
brown, let stand warm for an hour, then	No. 2
add the following dissolved by boiling.	Indelible Ink, Stable Against Water, Oil,
Water 200 g.	Alcohol, Alkali, Oxalic Acid, Chlorides
Borax 1.5 g.	a. Shellac 4 g.
Shellac 3 g.	Borax 2 g.
Aniline Blue 4 g.	Water 36 g. Boil till dissolved.
Name of the second of the other contents of the second of	
Non-Corrosive Writing Ink	b. Gum Arabic previously 2 g.
Gall Nuts 28 g.	Water \int dissolved 4 g.
Aniline Blue 6 g.	Mix a and b , boil, filter, cool, add
Ferrous Chloride 30 g.	c. Indigo Carmine to desired color
Glycerin 2 g.	Note: Just traces of sulphuric or
Hydrochlorie Acid 30 cc.	hydrochloric acids or salt make ink in-
Arsenic Acid 1 g. Phenol 1 g.	delible.
Phenol 1 g. Water 1000 l.	T 1 0 TT 11 0 0 17 7 17
-	Ink for Writing on Celluloid
Powdered Writing Inks	Ferric Chloride 10 g.
Formula No. 1	Tannic Acid 15 g.
Gallic Acid 10 g.	Acetone 100 cc.
Ferric Sulphate 10.7 g.	Dissolve the ferric chloride in a por-
Oxalic Acid 2 g.	tion of the acetone and the tannic acid in the remainder and mix the two. Use
Soluble Blue Dye 3.5 g.	any pen.
Formula No. 2	any pen.
Gallic Acid 10 g.	701 .1 7 .1 7 .1
Ferrous Sulphate Crystals 15 g.	Black India Ink
Tartarie Acid 1 g.	a. Borax 0.3 g.
Soluble Blue Dye 3.5 g.	Shellac, Wax-Free 1.5 g.
Indelible Inks	Water (Boiling Hot) 4 g. b. Black Tar Dye, Water-
	Soluble 0.1 g.
Formula No. 1	Water 4.1 g.
a. Chinese Gall Nuts,	Mix cold.
Powdered 750 g.	
Water, Hot 3000 g.	Non-Coagulating India Ink
Stir, keep standing 2 days, then press	
out extract; add to the extract:	Japanese Patent 110,282
b. Ferric Sulphate in Water,	Glue (Previously Heated at
(sp. gr. 1.48) 48 g. Solution, Saturated, of	120° C. for 3 hr.) 30 oz. Urea 10 oz.
Oxalic Acid 18 g.	Potassium Nitrate 60 oz.
10 g.	2 Companie Attorneo 00 02.

Urotropine	10	oz.
Carbon Black	60	oz.
Water	1000	07.

This ink will not coagulate at temperatures down to -30° C.

Silver Glow Ink

	VIII I CI	CION	TILL		
Tin				1	OZ.
Mercury				2	oz.

Grind together until liquid; then grind with 1 pint of 2% gum arabic solution. When used as an ink the writing will resemble silver.

Marking Ink for Chemical Porcelain Cobalt Oxide, Black Commercial Bismuth Subnitrate 1.2 g. Grind these together thoroughly with

Grind these together thoroughly with
Turpentine 15 cc.
Dresden Thick Oil 15 drops

Mark the porcelain with a pen, heat slowly to evaporate the liquids, and then ignite strongly. The porcelain apparatus is then ready for use.

Ink Erasing Fluid

An alkali hypochlorite, first applied to the ink to be removed; followed by an application of dilute acid, will remove ink from paper.

Ink for Glass or Polished Metal Sodium Silicate 2 oz. Liquid India Ink 10 oz. Use on clean surface with a steel pen.

Ink for Glass

Turpentine	20 g.
Venice Turpentine	6 g.
Shellac	10 g.
Mastic	2 g.
Lampblack	6 g.

The lampblack is added gradually to the mixture of other ingredients. This ink is very efficient for writing on glass photographic plates and lantern slides.

Stencil and Marking Ink U. S. Patent 2,002,939

Shellac Solution (4	lb. per	
gal.)	32	oz.
Turpentine	5.3-6	OZ.
Beeswax	2.0-2.3	oz.
Lampblack or Chr	ome	
Yellow	5.7-8	oz.
Alcohol	80-167	fl. 0"

Ink for Writing on Carbon Paper U. S. Patent 1,988,723

Titanium Dioxide	1	oz.
Mineral Oil	2	oz.
Mineral Spirits (Naphtha)	4	oz.

Carbon Paper Ink French Patent 774,922

Cottonseed Oil	1	lb.
Prussian Blue	1	lb.
Carnauba Wax	2	lb.
Paraffin Wax	2	lb.
Ozokerite	1/2	lb.
Octadecyl Alcohol	1	lb.

Transfer Ink

U. S. Patent 1,990,193

Carnauba Wax	3 lb.
Boiled Linseed Oil	2 lb.
Caustic Soda	0.375 lb.
Pigment	to suit

Thermographic Printing Ink U. S. Patent 1,992,016

Paracumarone Resin	100	lb.
Dibutyl Phthalate	50	lb.
Butyl Stearate	50	lb.
Drier	$2\frac{1}{2}$	lb.

Rotogravure Ink French Patent 776,825

Ethyl Cellulose	5	lb.
Alcohol	155	lb.
Alcohol Soluble Dye	40	lb.

Offset Printing Ink U. S. Patent 1,989,250

Pigment	34.4	lh.
Linseed Oil	21.5	
Varnish	33.2	lb.
Castor Oil	2.2	lb.
Stearin	3.7	lb.
Turpentine	5	lb.

Intaglio Printing Ink U. S. Patent 1,962,823

A pigment is used with the following binder:

	The state of the s			
	Rosin	2	lb.	
	Caustic Potash (10%)	1.6	lb.	
	Casein	0.1	lb.	
	Ammonia (sp. gr. 0.91)	0.24	lb.	
	Turpentine	0.2	lb.	
١.	Water	4	lb.	

INKS AND MARK	ING COMPOUNDS	191
Lithographic Bronze Printing Ink Varnish German Patent 604,019 Polymerized China Wood Oil 10 lb. Linseed Oil, Boiled 5 lb. Turpentine 2 lb. Carnauba Wax 1 lb.	Newspaper Ink Pit Coal Tar (0.85–0.89 Density) Linseed Oil Boiled with Litharge or Linseed Oil-Colophony Varnis	1 kg. 4 kg. sh 4 kg.
Polymerize China wood oil at 240-280° C., add linseed oil and heat to 200° C. for 2 to 3 hours. Cool and add the carnauba wax and turpentine. About 9 lb. of above is stirred with 16 to 18 parts bronze powder. Printing Lacquer	Pyroxylin Printing In Ethyl Oxalate Nitrocellulose (½ sec.) Dye (Basic) or Pigment	k 10 lb. 3 lb. 2 lb. 2 lb.
U. S. Patent 1,996,846 Nitrocellulose about 10 parts, ester gum about 25 parts, xylol about 30 parts, fenchone about 30 parts, dibutyl phthalate about 5 parts and pigment about 25 parts relative to the total of the other ingredients.	Typographic Ink Red Yacca Gum, Powder Borax Solution, Boiling Glycerin Gum Arabic Soluble Nigrosine Water	15 g. 4 g. 1 g. 2 g. 5 g. 73 g.
Solid Color for Rubber Printing Blocks Hansa Yellow 200 g. Alcoholic Shellac (50%) 50 g. Borax 50 g. Water 250 g.	Water-Soluble Printing Glycerin Gum Arabic Water Soluble Dye	Ink 100 oz. 50 oz. 10 oz.
Ink for Rotary Press Pit Coal Tar (Density 0.85-0.89) 100 g. Treat warm with: Sulphuric Acid (66° Bé.) 3 g. then neutralize with stirring by Soda Ash. Deodorize with calcium chloride	Lithographic Color In Glycerin Copaiba Balsam Venice Turpentine and Sandal Wood Oil Petroleum Oil Pine Turpentine	k 10 g. 20 g. 5 g. 2.5 g. 2.5 g.
and hydrochloric acid. Above Tar plus Pig Fat Glycerin To this liquefied and cleared varnish add Campêche Extract Campêche is described and cleared to obtain:	Alcohol Manganese Dioxide This mixture, prepared on bath, is thinned with Chloroform Ether Ammonia (28° Bé.)	5 g. 2.5 g.
Black, brown or violet coloration with Alum Copper Sulphate Potassium Bichromate Finally mix with Lamp Black 10 g.	Lithographic Ink for Repro Resin, Damar Petroleum Oil Glycerin Linseed Oil Varnish Color	

***************************************	-
Typographic Ink for	Newspapers
Colophony Tar	37 g.
Rosin Oil, Rectified	40 g.
Thinner: Petroleum	20 g.
Filter hot.	

T. marpre	Lamograpme	Ink	
Damar		50	oz.
Kerosene		100	oz.
Pigment		100	oz.

192 THE CHEMICA	L FORMULARY
Fine Lithographic Ink Asphalt (Gilsonite plus 60% of Rosin Oil plus '70 to 120% of Rectified Tar) 15 g. Pit Coal Tar 30 g. Paris Blue 2 g. Bone Black 3 g. Lamp Black 23 g. To get a typographic ink, increase the proportion of tar, and reduce sensibly the proportion of the color. Typographic Ink for Prints Colophony Tar 95 g. Rosin Oil (Medium, Neutral, Rectified) 50 g. Linseed Oil, Light 13 g.	Medium Varnish (for Inks) Rosin Oil 95 g. Crude Linseed Oil 35 g. Sulphonated Rosin Soap 7 g. Colophony 40 g. Evanescent (Invisible) Inks Formula No. 1 Cobalt Chloride 1 dr. Mucilage of Acacia 1 dr. Distilled Water 1 oz. Dissolve. The writing becomes blue when the paper is heated, and disappears again on cooling. No. 2 *Oxalomolybdic Acid 15 gr. Distilled Water 1 oz. Dissolve. Write with this in a dull
Lithographic Inks with Oil-Varnishes Thickened by a Resin Glycerin 40 g. Varnish, Medium 40 g. Soda Ash 2.8 g. Cream of Tartar 1.4 g. Venice Turpentine 16 g. Color 6-34 g. Tartar and soda are first dissolved in glycerin.	light. When exposed to sunshine, the writing appears blue; when wetted, the blue changes to black. * Made by dissolving Molybdic Acid to saturation in a hot solution of oxalic acid, and collecting the crystals on cooling. No. 3 Nickel Chloride 10 gr. Cobalt Chloride 10 gr. Distilled Water 1 oz. Dissolve. The writing becomes green
Varnish for Lithographic Inks Sandarac 15 kg Olive Oil 15 kg White Beeswax 12.5 kg Stearic Acid 12.5 kg Oleic Acid 2.5 kg Castile Soap 2.5 kg Burgundy Pitch 40 kg Stearin Pitch 10-20 kg Varnish for Artistic Prints Medium Strength Colophony, Pale 110 g Copaiba Balsam 70 g Tolu Balsam 2.5 g Benzoin Amygdaloid 3 g Linseed Oil 50 g Dissolve hot.	on heating. No. 4 Lead Acetate 10 dr. Distilled Water 1 oz. Dissolve. The writing is invisible, and becomes black when damped with a sulphide solution. Billiard Chalk Formula No. 1 Calcium Carbonate, Precipitated 115 g. Gypsum, Calcined 35 g. Pigment Powder (Blue, Green) 50 g. b. Borax Water (2%) about 180–200 g. to make a pasty liquid This paste is poured into slightly oiled molds. No. 2
Medium Varnish (for Inks) Rosin Oil 50 g. Sulphonated Rosin Oil Soap 3.5 g. Boiled Weak Linseed Oil 4 g. Boiled "Middle" Linseed Oil 52 g. Colophony 25 g. By removing the weak linseed oil, a strong varnish is obtained.	Calcium Carbonate 100 g. Gypsum 30 g. Borax Water (2%) 115–130 g. As above. Cellulose Tranfer Inks Formula No. 1 Cellulose Acetate 170 oz. Triacetin 200 oz.

INKS	AND MAR	KING C
High Phenol Resin Pigment No. 2	200 oz. 250 oz.	Hig
		Pet
Nitrocellulose (½ sec.) Triphenyl Phosphate Blown Castor Oil Basic Dye	15 oz. 20 oz. 5 oz. 2 oz.	Hig
Acetone No. 3	50 oz.	Xyl Peti
Nitrocellulose (½ sec.) Glyptal Balsam Stearic Acid Pigment Acetone No. 4	15 oz. 20 oz. 5 oz. 10 oz. 50 oz.	Ir Higl Tetr
Nitrocellulose (½ sec.) Phenol Formaldehyde Resin Beeswax Acetone No. 5	15 oz. 25 oz. 50 oz. 50 oz.	Use
Triphenyl Phosphate Butyl Tartrate Cellulose Acetate Mineral Oil Basic Dye No. 6	50 oz. 50 oz. 50 oz. 5 oz. 20 oz.	A su the ren contain about 3 of 95% of dieth
Ethyl Cellulose, High Viscosity Castor Oil Mineral Oil Bronze Powder Benzol	50 oz. 25 oz. 10 oz. 20 oz. 50 oz.	Potas Citric Mix t Water
Emulsifiable Transfer I	nk	
Diglycol Stearate	20 oz.	
Ethyl Cellulose	5 oz.	
Sodium Abietate	10 oz.	
Pigment	10 oz.	
7		Calciu Miner

Ink Remover

For cleaning dry printing ink from printers' rolls and type.

Denatured Alcohol Commercial Toluol Heavy Naphtha Creosote Oil	2½ gal. 1¼ gal. 3¾ qt. 1¼ gal.	
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Non-Inflammable Ink Remover (for Washing Printers' Rolls and Type) Carbon Tetrachloride 10 pt.

Carbon Tetrachloride 10 pt.
Toluol 13 pt.
Heavy Naphtha 11 pt.
Greosote 2 pt.

Printing Form Cleaner Use light gasoline.

Printing Roller	Cleaner		
High Test Benzine	90	ſl.	oz.
Petroleum	10	fl.	·0Z。

General Printing	Cleaner	
High Test Benzine Xylene	80 fl.	
Petroleum	15 fl.	
1 Coloredin	5 fl.	oz.

Intaglio	Printing	\mathbf{Press}	Cle	$an\epsilon$	er
High Test : Tetralin	Benzine		80 20		
			20	11.	UZ.

Off-Set Printing Cleaner Use light petrol (gasoline).

Ink Remover U. S. Patent 1,968,304

A substantially non-aqueous cream for the removal of ink stains from the skin containing about 500 g. of zinc stearate, about 300 g. of citric acid, about 500 cc. of 95% ethyl alcohol and about 2000 cc. of diethylene glycol.

Ink Eradicator		
Potassium Alum Citric Acid		lb.
Mix thoroughly and dissolve in	_	
Water	3	lb.

Stencil Coating Paste U. S. Patent 2,011,898

Formula No. 1 Calcium Oleate Solution

Calcium Oleate 20 oz. Mineral Spirits 80 oz.

The above ingredients are combined by heating for a short time in a steam-jacket kettle.

No. 2

Ammonium Stearate Solution Ammonium Hydroxide

(sp. gr. 0.9) 0.41 oz. Water 98.84 oz. Stearic Acid 0.75 oz.

The stearic acid is broken up into small pieces and agitated with the other ingredients until dissolved.

No. 3

Ammonium Oleate Solution

(sp. gr. 0.9) 0.41 oz. Water 98.84 oz. Oleic Acid 0.75 oz. The above ingredients are combined in

the same way as those of No. 2.

Suitable compositions for stencil paste in which the false bodying agents are incorporated are given below. The composition of the particular resin used is given after the examples setting forth the stencil paste compositions.

No. 4

White Stencil Paste

Lithopone	46.1	oz.
Zinc Öxide	23.1	
*Resin A	15.7	0Z.
Drier	1.5	oz.
Ammonium Stearate Solu-		
tion of No. 2	2.3	oz.
Calcium Oleate Solution		
of No. 1	4.8	
Mineral Spirits	6.5	oz.

No. 5

The same composition as No. 4 except that 23 parts of the lithopone are replaced by 23 parts of diatomaceous earth. The effect of the soap solutions described in the preceding examples is enhanced by the use of cellular or fibrous materials such as diatomaceous earth or "Asbestine."

No. 6

The same composition as No. 4 except that basic lead carbonate is substituted for lithopone.

No. 7

The same composition as No. 4 except that resin B is used instead of resin A.

No 8

Black Stencil Paste

Carbon Black	17	oz.
"Asbestine"	4.1	oz.
†Resin B	64	oz.
Drier	4.1	oz.
Ammonium Oleate Solu- tion of No. 3 Calcium Oleate Solution	7.2	oz.
of No. 1	3.6	oz.

No. 9

The same composition as No. 8 except that 7.2 parts of ammonium oleate solution are replaced by 4 parts of mineral spirits and 3.2 parts of calcium oleate solution of No. 1.

No. 10

Red Stencil Paste

Toluidine Red	19.8	
Barytes *Resin A	28.6	oz.
*Resin A	21.8	oz.
Ammonium Stearate Solu-		
tion of No. 2	15.4	oz.
Mineral Spirits	12.8	oz.
Drier	1.6	oz.

The linseed oil modified resin given in this formula may, if desired, be replaced by a resin modified by linseed oil acids such as indicated by resin C below.

The ingredients in the pastes described above are combined in accordance with the usual products of paint manufac-

The following resins are illustrative of the class of polyhydric alcohol-polybasic acid resins especially suitable for the purposes of the present invention. These resins are made in the conventional way by reacting the ingredients in the proportions indicated.

*Resin A Glycerol Phthalic Anhydride Linseed Oil	12.8 oz. 28 oz. 59.2 oz.
	59.2 OZ.
†Resin B	
Glycerol	15 oz.
Phthalic Anhydride	35 oz.
Linseed Oil	50 oz.
Resin C	
Glycerol	17.1 oz.
Phthalic Anhydride	27.1 oz.
Linseed Oil Acids	55.8 oz

LEATHER, SKINS, FURS

Chamois Leather from Rejected Calf Skins

The skins are soaked, pasted with sodium sulphide 1 and calcium oxide 4 (25° Bé.) at a temperature not exceeding 30° C., limed with calcium oxide 10 g. per liter, sedium sulphide 4 g. per liter, water 400%, at 20° during 18-20 hours, washed with water at 22° for 40 minutes, fleshed, treated with 0.3% hydrochloric acid and 2% sodium chloride (of the weight of the raw skins) at 25°, softened with a concentrated softener (0.1% of the weight of the raw hide), for 1 hour at 35-37°, pickled for 40 minutes with hydrochloric acid 1.7, sodium chloride 7 and water 80%, tanned with chrome extract of 2% chromic oxide, having a basicity of 50%, split, neutralized, washed, greased, with 0.5% alizarin oil, 2% egg yolk and 150% water, washed with water at 35°, dried at 35°, let stand 2 days, dehaired in sawdust, stretched, cut, sand-papered and soaked.

Chamois Leather of Natural Color from Rejected Kid Skins

The skins are soaked in water at 18-20° C., drummed for 45 minutes at 17° fleshed, soaked again in water at 16-17°. drained and treated with a mixture of sodium sulphide 2%, calcium oxide 5% (of the soaked skins) of 30° Bé. at a temperature of 35-40°. The hair is removed by hand and the skins are placed in a lime solution for 5 days at 12-16°. They are then washed for 30 minutes, split, the thin parts are tanned by the formalin-fatty method and the heavier parts are chrome-tanned. The flesh side is treated with 0.5% hydrochloric acid for 45 minutes. The skins are further pressed and drummed in 5% of seal fat, and treated in an oxidizing chamber for ½ hour at 32-35°. The above processes are repeated except that the oxidizing drying is carried out at 40-42°. The product is stored for 3 days, degreased with 200% water at 45° and sodium carbonate solution (5% of the weight of the skins) is added, the liquid discharged and the above soda solution again added together with water. The goods are soaked with water at 40-45°, drained, dried and stretched. They are dyed with nigrosine, drummed for 6-7 minutes and fat liquored with 0.75% castor oil, 2% alizarin soap and 2% rosin soap.

Velure from Rejected Pig Skins

Soak the raw hide in pieces weighing 1-3.5 kg. to a liquid factor of 1.5, at 20° C. and for 2 hours treat with 1 part sodium sulphide and 3 parts calcium oxide, density 25° Bé., at 25° let stand for 3-6 hours, unhair, wash and sort. Then treat with sodium sulphide 10 g. and calcium oxide 10 g. per liter at 20° for 4 days, split to an average thickness of 1.25 mm. and wash to a liquid factor 1:5 at 20° for 2 hours. Treat with concentrated softener 0.5% at 37° for 2-3 hours. de-ash with bisulphite 2% at 28° at a liquid factor 1:5; wash to a liquid factor of 1:5 at 25° and during 30 minutes. Pickle with sulphuric acid 2%, sodium chloride 10% and water 80% for 40-60 minutes at 18°. Tan with chrome extract containing chromic oxide 1.8, basicity 45 and water 80%; to complete tanning the basicity may be raised if necessary. Neutralize with bicarbonate 1.25 and water 200% at 35°, wash with water 300% at 40° for 30 minutes. Fat liquor with alizarin oil 1, egg yolk 3, water 150% at 40° for 40 minutes, and wash with water 300% at 35° for 25 minutes. Dry at 35°, unhair in sawdust containing 60% water for 16-20 hours, stretch, cut and polish.

Chrome-Tanned Black Calf-Leather Chamois

A calf leather which was previously tanned is planed on the grain side, neutralized, treated with 2% of pure fats, dried, unhaired and nailed on frames. The skin is then worked over with grinding stones and the final treatment is given with pumice stone. Skins with a light nap are worked over with a wire brush (by hand). The skins are finally dyed with 15% (of their dry wt.) of substantive dyes and 4.5% ammonia, the mixture being diluted with 50% water.

Preparing Leather from the Mucous Stomach Membrane of Cattle

(1) The material is soaked, slightly fleshed, limed for 2 days, with about 12% slaked lime on the weight of the tissue, washed and delimed with bisulphite. Tanning by vegetable or by onevat or two-vat chrome methods is followed by the usual dyeing, fat liquoring, drying and finishing. (2) The membrane is soaked for 2 hours in cold water, then for 15-20 minutes each in 3 vats with a gradually increasing temperature from 22° C.

Removing Scales from Shark Skins

Give the skins a salting in a 1% solution of sodium chloride. Then a treatment in a ½% solution of hydrochloric acid. This method should dissolve the scales, but if for any reason it does not, keep on increasing the percentages of both materials. Then give the skins a thorough washing in pure water in a drum. Watch carefully that the hydrochloric acid does not attack the skins themselves.

Loosening Hair from Hides Canadian Patent 353,326

Wheat Sho	orts		14	lb.
Wheat Bra			6	lb.
Phenol Sol	ution	$(2\frac{1}{2}\%)$	0.6	cc.
Water			15	gal.

Preparing Pigskins for Tanning

First, scrape the raw skins until they are nearly dry. Then give them a good soaking for a day or two. Next wash them in a drum or vat containing a warm solution of sal soda or similar product for loosening the grease. In preparing this solution, use from 1% to 2% of sal soda according to the condition of the skins, i.e., they appear to be extremely greasy, a higher percentage of sal soda is preferred. After the skins have received a thorough soaking in this solution, strike them out thoroughly with a dull knife, forcing out as much grease as possible. Very greasy skins should be struck out two or three times. Then rinse them off in warm water and soak them overnight in cold water, after which they are unhaired and limed.

As pigskins absorb tan liquors somewhat slower than calf and other skins, it is good practice either to give them slightly stronger liquors or a longer time in the same strength liquors you are using for your other stock. This sugges-

tion applies more especially to a vegetable tannage.

Pigskins being of a very greasy nature require less oiling or fat liquoring than other skins. Some tanners reduce the oiling from 20% to 30%.

Felting Animal Hair German Patent 608,770

Hair is rendered capable of fulling and felting by treatment with a bath containing small amounts of oxy acids of metals of the chromium group or their salts together with hypochlorous acid or persulphuric acid or their salts. Thus pelts are treated with an aqueous solution containing 2% potassium chlorate, 1% nitric acid and 0.1% chromium in the form of dichromate at 10–100° C., and dried.

Treating Lizard Skins

Bleaching should be effected in two solutions. (1) potassium permanganate 5 g. per liter, sulphuric acid 1 g., water 500% of the weight of the skins, and (2) water 500%, bisulphite 25 g. per liter. The washed skins are dyed beige by treating with 0.03% orange PB, 0.04% methanyl yellow and 200% water for 20 minutes, adding 0.3% acetic acid and treating 20 minutes. For gray use nigrosine 0.1%, acid brown 0.01% and water 200% at 45° for 15 minutes; add 0.3% acetic acid and treat for 15 minutes. For violet use wool brown 0.5% and acetic acid 0.5% at 45°, add 0.1% methyl violet after 30 minutes and treat for 15 minutes. For blue use sulphone acid-blue 0.3% and water 200% at 45° C. for 15 minutes, add 15% acetic acid and treat for 20 minutes.

Bleaching Deer Skin

Formula No. 1

Make a bath with

Hydrogen Peroxide (30%) 5-8 lb. Seignette Salt 0.5 lb. and put the skins into it for ½ hou

and put the skins into it for ½ hour. Dry them thereafter at 30° C. If the skins are not pale enough, repeat in the same bath.

No. 2

Put skins into a solution of Potassium Permanganate

Potassium Permanganate 3 lb. Sulphuric Acid 0.5 lb. Water 96.5 lb.

for 30 minutes, moving repeatedly.

Wash out in cold water, then in solutions of

Sodium Bisulphite Powder 95 lb. Water 95 lb. (for ½ a minute), and Hydrochloric Acid 5 lb. Water 95 lb.

(for ½ minute).

Then wash out very carefully, repeat the process until the wanted paleness is reached.

No. 3

Tanning After Bleaching (Often Advisable)

Wash for 2 hours at 30-35° C. in solution of sodium carbonate, spill with water, and treat for 7 hours in a solution of

Sodium Carbonate 2 lb. Formaldehyde (40%) 2.5 lb. Water 95.5 lb.

Tanning Greenland Seal Skins

The sorted skins are soaked in water for 10 minutes, fat is removed from the flesh side and the skins are again soaked in water for 36 hours with change of water at 12-hour intervals. They are degreased in a drum charged with water of 30° C. with addition of 1% sodium hydroxide (calcined on the salted skins). The skins are washed in running water for 30 minutes, drained on racks for 2 hours, placed for 30 minutes at 25-30° in a solution prepared from sodium sulphide 20 g. per liter and calcium hydroxide 160 g. per liter, unhaired with a tool, washed till the concentration of sodium sulphide amounts to 20-25 g. per liter, and treated for 2 days in a lime solution used once for unhairing, with addition to the solution of 12 g. calcium hydroxide in the course of the processing. The skins are then washed, split, delimed and tanned in a six-vat battery for 6 days, with a 4° Bé. solution in the first and 4.25° Bé. in the last vat. The drum tanning may be carried out in an oak extract of 7° Bé. The aging in stacks requires 24 hours and the deacidification, which is carried out with 1° Bé. solution during 4 hours, is followed by washing in running water for 8-10 hours.

Tanning Horsehide
Full Grain Horse for Glove and Sport
Goods

Having selected hides after unhairing for this type of leather, they are pickled, tanned, pressed, staked, etc., in the same manner as buffed glove horse. The stock is then split and shaved. After this it is neutralized and fat liquored in the same manner as for the "One Bath" tanned stock which is given below.

One Bath Tannage for Full Grain Horse

Often a tanner prefers to tan glove horse leather with the single bath tannage rather than with the two bath tannage. The final results will be the same, as both tannages produce excellent leather.

After lime splitting, the stock is bated and washed and taken to the chrome tan wheels. Maximum loads of 3000 lb. of lime split stock are considered sufficient. The tannage is based on this weight.

Place the stock into the drum with 180 gal. of water and 180 lb. of salt, mill for 10 minutes, then add 45 lb. 66° Bé. sulphuric acid in 15 gal. of water. Mill for 75 minutes, then add 42 gal. of chrome liquor.* This is added in three doses of 14 gal. each, 30 minutes apart. After the last addition is made, continue milling for 4 hours, let stand in drum over night.

night.
The following morning mill the stock

1/2 hour, then add:

Fifteen pounds bicarbonate of soda, first dissolved in 20 gal. of water. Add this at the rate of 1 gal. every 2 minutes; continue milling 30 minutes, remove from the drum, lay flat on trucks, let drain for 24 hours, set out, split and shave.

*The chrome liquor used for this purpose is made as follows:

 Bichromate of Soda
 1000 lb.

 Aluminum Sulphate
 400 lb.

 Sulphuric Acid, 66° Bé.
 800 lb.

 Corn Sugar
 250 lb.

 Total Volume
 500 gal.

Use a lead lined tank. Place the bichromate, aluminum sulphate and 200 gallons of water into the tank, agitate well by means of an air line, then add the sulphuric acid. The corn sugar is made to a syrup with water and is added very slowly, taking the usual precautions.

After all the sugar has been added, add five gallons of bisulphite of soda, 33° Baumé, boil the liquor for one-half hour, allow to cool and make up to 500 gallons, stir well and allow to age ten days before using.

Coloring

Divide the tanned split stock into lots of 400 lb. each for coloring and fat liquoring. Place the stock into the drum with 120 gal. of water at 90° F., then add 6 lb. of bicarbonate of soda dissolved in 20 gal. of water, and mill for ½ hour. Drain the drum and wash the stock for 1 hour at 80° F., again drain

the drum and add 200 gal. of water at 120° F.

Prepare the following dye mixture:

Boil together in 30 gal. of water, cool to 125° F., and add to the drum. Mill stock in the dye solution for ½ hour, then drain the drum.

This will produce a cream color which is a standard for glove and sport goods stocks. The amount and type of fat liquor determine the purpose to which the stock will be used.

Fat Liquor for Stretchy Glove Leather
Sulphonated Cod Oil 24 lb.
Sulphonated Mineral Oil 24 lb.
Sod Oil 24 lb.
Borax 4 lb.

Place the materials into a barrel in the order given, stirring well upon addition of each item. Add 25 gal. water and heat by means of a steam jet agitator to 160° F. The total volume should be 50 gal. This emulsion is added to the drum after draining off the exhausted dye solution. Mill for one hour, rinse very slightly with water at 100° F., take out of drum and horse up for 24 hours, then hang up to dry.

Fat Liquor for Sporting Goods Leather
Sulphonated Mineral Oil 64 lb.
Sod Oil 24 lb.
Borax 4 lb.

Place the materials into a barrel in the order given, stirring well. Then add 25 gal. of water and heat by means of a steam jet agitator to 160° F. The total volume should be 50 gal. This emulsion is added to the drum after draining off the exhausted dye solution. Mill for 1 hour, rinse slightly with water at 100° F., take out of drum, horse up 24 hours, then hang up to dry.

Drying

This type of leather can be dried rapidly. Since it is quite wet, the initial air temperature can be 120 to 130° F. Rapid circulation of the air must accompany the high temperature; the moist atmosphere is gradually expelled from the dry room, emitting at the same time fresh air and reducing the temperature, so that the stock is thoroughly dried in 24 hours.

Crusted Stock

After the stock is dry it is crusted for five days. Dip the crusted stock in

water at 110° F. for one minute, place into bins, cover well with damp burlap and allow to mull for four hours. Then place into damp sawdust (containing about 35% moisture) and let it rest for 24 hours. Then stake on a Slocum Machine and hang up to air off for an hour.

Dry Mill

After the stock has aired, place into a dry mill. For each 100 sides use 10 to 20 lb. of French chalk, the amount depending upon the size of the stock. Dry mill for 1 hour. Remove from dry mill and stake on the Baker Machine. After the second staking, polish the grain on a shearling wheel.

Notes: Some adjustments may have to be made for either the "one bath" or the "two bath" operations. In a greater number of cases the adjustment is made in the fat liquor stage, either increasing or decreasing the amount. Drying of the stock must be carefully controlled since this operation is very important to a soft, yet full feeling leather.

Leather of this type should not be tacked. Leather of this type should be stretchy, the glove more so than the sport leather. The latter is used principally for baseball gloves.

Black Garment Horse Leather

This type of leather is used principally for coat stock, although it can also be used for glove purposes. The market for this leather is highly competitive and therefore the leather must be made as economically as possible. Sheep, in grain and suede, is used very extensively and is produced at a low cost including the raw material. Because of this, it has found a greater market than horse leather. For general utility and durability, horse garment leather excels sheep leather.

The stock is sorted in the beamhouse before bating. The butts should be split down to a minimum. After bating and washing, the stock is transferred to the chrome tan yard.

A maximum drumload of 3000 lb. of lime split stock will be used. The stock is placed into the drum with 200 gal. of water at 65° F. and 180 lb. of salt. Mill 5 minutes and then add 42 lb. sulphuric acid, 66° Bé., in 15 gal. of water, and mill 15 minutes, then add 45 gal. of chrome liquor.* This is added in three doses of 15 gal. each, 30 minutes apart. After the last addition of chrome liquor mill for 5 hours, let stand in drum overnight.

The following morning, mill the stock for 30 minutes, then add 15 lb. of bicarbonate of soda dissolved in 20 gal. of water at 75° F.

Add the soda at the rate of 1 gal. every 2 minutes. After the last addition mill the stock for 30 minutes, remove from drum and horse up for 24 hours, set and split. The split stock is divided into lots of 500 lb. each for coloring and fat liquoring.

*The chrome liquor for this tannage is made as follows:

Bichromate of Soda	1000 lb.
Sulphuric Acid, 66° Bé.	980 lb.
Corn Sugar	332 lb.
Total Volume	500 gal.

The usual precautions must be taken and the manner for procedure is the same as that for the chrome liquor under "One Bath Tannage" for glove horse.

Coloring

Place the stock into the drum with 150 gal. of water at 90° F., and add 3¾ lb. soda ash in 10 gal. of water. Mill for 30 minutes, then wash with water at 110° F. for 1 hour. Drain the drum and add:

Water at 120° F. 250 gal. Direct Black in 30 gal. of Water at 120° F. 17½ lb.

Mill 30 minutes and drain the drum, add:

Water at 120° Methyl Violet	F.	$250 \\ 2\frac{1}{2}$	
and			

Acetic Acid 4 oz. in 20 gal. of water, mill 20 minutes. Drain drum.

Fat Liquoring

Prepare the following:

Logwood Crystals	$7\frac{1}{2}$	lb.
Water, Boil and Add	20	gal.
Fig Soap	15	lb.
Sod Oil	100	lb.
Sulphonated Cod Oil	10	lb.
Total Volume	50	gal.

Use steam jet agitator for the purpose of preparing the above emulsion, add to the drum at 150° F., and mill 1 hour. Remove from drum and horse up to drain for 16 to 24 hours, set out on Turner Serial Table Machine.

Oiling and Drying

Oil off the set out stock on the grain with a light paraffin oil, using a shearling swab for the purpose. Apply a light coat. Then send the stock to the dry room. Hang up the stock in a room equipped with fans and heating coils. A temperature of from 90 to 100° F. is maintained; the air is well circulated with fans so that drying is effected in

24 to 36 hours. The stock is then crusted for two days in a cool room.

Sammying and Staking

Dip the stock in warm water, 110° F., for 1 minute, place into a bin, cover with burlap and allow to mull for 4 hours, then place into damp sawdust containing 40% moisture, let rest for 24 hours. Then stake on a Slocum Machine equipped with a fiber pad on the staking head. Apply as much pressure as the stock will stand; cracking of the grain must be avoided. Then hang up the stock to air off at room temperature, restake and trim closely where necessary and again stake if hard spots are found.

Finishing

Use the following finish:

Shellac Solution	6 pt.
Casein Solution	8½ pt.
Liquefied Gelatin	6½ pt.
Carnauba Wax Emulsion	1½ pt.
Sulphonated Cod Oil	1 pt.
Nigrosine	1¼ lb.
Water	30 pt.
Ammonia	1 pt.

Mix the above ingredients in the order given, the Nigrosine first dissolved in the water. Apply two coats of the finish to the stock, allowing to dry well after each application. Finally polish on a shearling wheel.

In order to obtain the desired results it may be necessary to vary the quantities of some of the finish materials. A third coat of finish may also be required. Proper drying between coats is of im-

portance.

The greatest factor affecting finishing of leather is the type and amount of fatliquor used. This holds particularly when a finish job at low cost is desired. In other words, the finish must be properly adjusted by varying its components until the proper balance is obtained.

Synthetic Tanning Process U. S. Patent 1,975,616

The hides, skins or pelts are prepared by any suitable and well known process and then immersed in a solution containing approximately 20% of a urea-formal-dehyde solution and 10% of salt at about 35° C. and gently agitated for about 5 hours. The temperature may then be raised to 45° C. and the solution acidified to about pH_3 with sulphuric acid and agitation continued for 30 minutes. The temperature is then raised to 55° C., the skins worked for 15 minutes, cooled, rinsed in cold water, neutralized with

sodium bicarbonate, rinsed, fat liquored and dried.

One method of producing the ureaformaldehyde solution mentioned in the above example is as follows: 3 oz. of urea, 11/2 oz. formaldehyde, 2 oz. sodium carbonate and 16 oz. of sodium chloride are dissolved per gallon of water, and this solution employed in the tanning process at once, or at least prior to the formation of an insoluble precipitate.

Leather Oil

Spindle Oil Caoutchouc, Crude	96 g. 3 g.
Resin, Coumarin, Viscous,	o g.
Liquid	1 g.

Heat to 100° C. and stir until dissolved; add a little Birch Tar Oil (as perfume).

Sport Leather Oil

Pale Train Oil	5 0	g.
Degras	20	g.
Woolfat, Neutral	5	g.
Birch Tar Oil	5	g.
Spindle Oil, Refined	20	ğ.
Melt together and add:		
Caoutchoue Solution (5-10%))	
in Toluol		g.

Oiling Leather with Petrolatum

Satisfactory penetration is obtained by drumming with a hot mixture of petrolatum 45, mineral oil 40, and degras 15%.

Special Leather Oil

Cold Test (20°) Neats-	
foot Oil	50 gal.
Paraffin Oil (28°)	25 gal.
Water	25 gal.
Sulphonated Castor Oil	J
(50%)	25 lb.

Manipulation: Mix water first with the sulphonated castor oil. Then mix all ingredients at 30° C.

Leather Fot Black

Deamer	1000	Diack		
Formula	No. 1	No. 2	No.	3
Paraffin Wax	8	4	5.2	g.
Wool Fat, Raw	2	1		g.
Montan Wax,				Ŭ
Crude	4	3	3.9	g.
Carnauba Wax,				0
Gray	2			g.
Nigrosine, Fat-		,		0
Soluble	1	0.3	0.39	g.
Train Oil	8	4	5.4	g.
Mineral Oil	60		32	g.

Leather Fat, Yellow

Formula	No. 1	No. 2	2
Paraffin Wax	8.5	5	kg.
Beeswax, Yellow	1.5		kg.
Train Oil	7	4	kg.
Spindle Oil	45-48	27	kg.
Yellow 1435, Dye	10	10	kg.
Carnauba Wax	-	1	kg.
Wool Fat		0.3	kg.
	_		
NT.	2		

No. 3

Paraffin Wax	8,000	
Carnauba Wax	1,375	g.
Wool Fat	340	g.
Train Oil	5,670	g,
Mineral Oil	35,000	ğ.
Yellow 1435, Dye	12	ğ.
		_

No 4

710° T		
Paraffin	10,000	g.
Ceresin	9,000	g.
Carnauba Wax Arrears		g.
Train Oil	4,000	g.
Spindle Oil	70,-80,000	g.
Yellow 1435	20	ø.

Leather Dressings or Finishes

Formula No. 1

Shellac Venetian Turpentine Castor Oil Alcohol	9 1 1 89	න් නේ නේ ප
Mix until dissolved and filter.		ş.

No. 2

Gum Mastic	12 g
Gum Sandarac	5 g.
Castor Oil	2 g.
Alcohol	81 g
No 2	8

Orange Shellac 16 Caustic Soda 0.9 g. 1.2 ğ. Boric Acid Sodium Ricinoleate 0.9 g. Water 81

No. 4

Orange Shellac	27 g.
Caustic Soda	21/4 g.
Boric Acid	$2\frac{1}{4}$ g.
Sodium Ricinoleate	$2\frac{1}{4}$ g.
Water	66¼ g.
3.T ==	

No. 5

Shellac, Bleached	20	g
Galipot Resin	1/2	g.
Borax	4	g
Ammonium Hydroxide	1/2	g
Turkey Red Oil	2	g.
Water	79	- 2

Dressing for Hunting Leather (Inflammable!)

a. Nigrosine Base	10 g.
Olein	30 cc.
b. Benzol (90%) Alcohol Cleaning Benzoline	150 cc. 500 cc. 500 cc.

Auto Top and Artificial Leather	Dressing
Nitrocellulose (Film Scrap)	40 g.
Camphor	10 g.
Ethylacetanilide	10 g.
Castor Oil	5 g.
Lampblack	5 g.
Nigrosine	2 g.
Alcohol	100 g.
Benzol	100 g.

Suede and Chamois Leather Dressing U. S. Patent 2,015,943

Acetone	90 oz.
Chloroform	60 oz.
Liquid Petrolatum	140 oz.
Naphtha	870 oz.

Leather Finishes

A good polish is made from 22 g. stearin, 22 g. carnauba wax and 56 g. inseed oil. It is better to prepare an "emulsion polish" by mixing 22 g. stearin, 22 g. carnauba, 11 g. paraffin, 23 g. linseed oil, 3 g. ammonium chloride and 17 g. water. The carnauba wax may be replaced by synthetic waxes. Waterproof spirit finishes are made by mixing shellac (9 g.), Venetian turpentine (I g.), castor oil (1 g.), and 96% alcohol (89 g.); or mastic (12 g.), sandarac (5 g.), castor oil (2 g.), and spirit (81 g.). All grease should be removed from the leather before application of spirit fin-For making polishes of good elasticity a recipe recommended is: ruby shellac (16 g.), technical caustic potash (0.9 g.), boric acid (1.2 g.), castor oil soap (0.9 g.), water (81 g.). Camphor oil may be added as a perfume. For treating leather of more porous nature, colloidal matter such as carragheen moss, algin, etc., are added to the above soap finishes, or gum tragacanth may be used. A recipe for green bronze finish is magenta 7.6 g., safranine 1.9 g., ruby shellac 1.4 g., and methanol 89.1 g.

Fur Glazing

Dissolve 3 to 6 oz. of paraffin wax in 1 gal. petroleum cleaning solvent.

Approved cleaning solvent is preferable because of its safety during ordinary handling.

Precaution: Paraffin separates from the petroleum solvent at temperatures below 70° F. At —15° F. it is completely

chilled out of the solvent.

This finish is used for the saturation of dry cleaned furs to replace any oils removed and to make them water repellent. It is also sponged or sprayed on materials that are lifeless or lusterless after cleaning and drying to produce high gloss.

Natural Color and Glaze for Snakeskins

An alum tannage is good for pocket-book leather and will as a rule impart a natural color. For each 100 lb. of bated and drained skins use 7 lb. alum, 2 lb. salt, 8 lb. flour and 5 lb. liquid egg yolk. The alum and salt are first dissolved in a small quantity of hot water and the solution then cooled. After cooling, the solution is added to the flour with constant stirring. Dissolve the egg yolk separately in a small quantity of cool water and then add to the other ingredients. This mixture when ready to apply should weigh about the same as the skins, that is, it should measure about 10 gal. for every 100 lb. of skins.

Stir the skins in this mixture for about 3 hours, or until nearly all of it is absorbed by the skins. Leave the skins in the same container or vat overnight. Then strike them out, stretch moderately on boards and dry. After drying, take the skins off the boards and wash them with a brush in cool water. This washing will remove any dried mixture re-

maining on the grain side.

Next lay the skins in piles overnight with grain to grain and cover with a moist cloth. Then stake and dry. After drying give the skins another staking. Some tanners also fluff the flesh side.

A good mixture for glazing can be made from the following: 1 oz. egg albumen, ½0 oz. gelatin, 2 oz. milk and 5 pt. water. The egg albumen is dissolved in 4 pt. of the water at 90-95° F. and the gelatin dissolved in 1 pt. of hot water and then allowed to cool to 90-95° F. The two solutions are mixed and then the milk is added.

This mixture is brushed on the grain side. The skins are then dried again and glazed by machine. Some tanners repeat this application and add a small quantity of casein or shellac. Others use

castor oil and methylic alcohol.

Dressing Bagdad Leather

Skins known commercially as Bagdads differ considerably in weight, size and quality, but they are all usually heavily loaded with dirt and loose tanning matters, all of which require to be completely removed before the goods can be properly dressed. After sorting, trimming and perhaps necking on the shaving machine, the goods need drumming for half an hour in a solution made up of 10% salt and 1/2% sulphuric acid on the dry weight. Some tanners use a cold solution, but a temperature of 100° F. will be found advisable for complete action. The object of processing the goods in the above liquor is to cleanse and open the pores of the leather so that it will be able to absorb the tannins during the next stage of dressing. At the end of the allotted time, namely half an hour, the liquor should be run off and the goods washed up in running water, preferably warm, for three-quarters of an hour. If the Bagdads are in a filthy condition the percentage of sulphuric acid should be increased to 1%, and this will generally prove strong enough to clear the grain and remove any stains, particularly iron marks. These preliminary processes are very important, especially in the case of whites, where it is of the utmost importance that the leather should be as clean as possible before the bleaching or whitening process commences.

Re-tanning

This operation can be successfully carried out in the drum, and, indeed, this is really the most suitable receptacle. A good synthetic tanning material, such as Maxyntan or Sellatan, in conjunction with sumac extract, usually forms the basis for a white tannage, and it is not advisable to use any tannin likely to darken the color of the leather. A run for half an hour in 5% of the synthetic followed by half an hour in 5% sumac extract will be found eminently satisfactory, but if it is necessary to reduce expenses to a minimum, the synthetic can be increased and less sumac extract employed. The tannage gives a very clean and fairly soft leather which will feed up well. The amount of water used depends a great deal on the weight and size of the goods, but in all cases the minimum should be run in, as this will ensure better exhaustion of the liquor.

After re-tanning for one hour, the goods should be taken out of the drum and horsed up overnight. Whilst this is not absolutely necessary, it is always ad-

visable if time and labor charges will permit, as it enables the tan to fix and the fibers to feed. Practical experiments have shown that there is a recognizable difference in the handle of leather allowed to drain for 12 hours as compared with leather rushed through the processes.

Bleaching

Next day, run the goods in the following solution: 2½% barium chloride and just sufficient warm water, 100° F., to cover the leather.

A run of a quarter of an hour will enable the leather to take up the barium salt and exhaust the solution. An addition of sodium sulphate, 5%, dissolved in a small volume of warm water will precipitate barium sulphate, a white insoluble salt, in the fibers of the leather. This bleaching process is quite economical and if worked properly it will be found to give a very clean, white leather.

Some tanners use sulphuric acid instead of the sodium salt, but sodium sulphate is equally satisfactory and with it there is less chance of the leather being

rendered hard and brittle.

Whitening and Filling

To fill out the leather, improve its handle and general appearance, it is advisable to work the goods in the following mixture, which should be added to the drum through the hollow axle:

Devolite Clay	15	lb.
Flour	15	lb.
Soap	5	lb.
French Chalk	5	lb.
Turkey Red Oil	21/2	lb.
Trace of Methyl Violet.		

A run of three-quarters of an hour in this liquor will complete the operation and afterwards the goods should be horsed up for a few hours preparatory to striking out and straining. The former process must be well done in order to remove all the wrinkles and drawn grain. To retain the fullness and suppleness of a well-nourished leather, the latter should be dried out in a moderate temperature. It is a bad practice to dry the leather in a fierce temperature for the sake of a few hours, but if this is imperative, then the temperature should be increased gradually. When dry, the leather requires buffing, then chalking on the grain and flesh, and finally boarding.

Semi-chrome Colors

A better quality skin is usually chosen for this work, and naturally the tanner has a better chance of producing a full and nice feeling leather. Goods should be washed in warm water for half an hour to remove loose dirt, and then stripped in a weak alkaline bath made up with 1 to 2% borax calculated on the dry weight of the leather. The stripping should take about an hour, and by this time practically all the loose tannin will be removed. The alkaline liquor should then be run off and the goods thoroughly washed in running water for half an hour.

Re-tanning in a Chrome Bath

After draining, the washed leather should be drummed with its own weight of a 4% salt solution for 10 minutes and the chrome liquor added. Prepare the chrome liquor by adding soda crystals to reduce the basicity. When using panchrome, 1 lb. of soda crystals for every 8 lb. of chromium salt is recommended. The latter should be dissolved in a known volume of hot water, and the soda dissolved in a small amount of hot water. The alkali must be added very slowly and the liquor stirred constantly during the addition.

The amount recommended for retanning Bagdads is 7% chromium salt on the dry weight of the leather. chrome liquor should be passed into the drum through the hollow axle in three parts, at intervals of half an hour. A period of 21/2 to 3 hours is recommended for complete re-tannage. The addition of 1% ordinary washing soda is then made, and drumming continued for a further At the end of that time, the leather should be well tanned, and it is advisable to horse up for twelve hours or The next morning, the goods will need neutralizing, and 1% borax on the dry weight is recommended; a period of three-quarters of an hour will be found to be sufficient to neutralize the leather.

A light mordanting is recommended to ensure more level dyeing, and to give the leather a better feel or handle. Gambier is quite good, so also is Osage Orange Extract; about 2% on the dry weight will be found ample. Acid dyes should be used and there is, of course, an unlimited number of colors available.

After dyeing, the leather should be well fat liquored, and the following recipe is excellent for semi-chrome clothing leathers. Dissolve ½ oz. of potassium carbonate in a small quantity of hot water, 180° F., and then add 2 lb. of neatsfoot oil and ½ lb. of potash soap. Emulsify the mixture and then add 1 lb. of heavy sulphonated oil and ½ lb. of mineral oil and stir vigorously until the emulsion is stable. Use 4 lb. of this

fatty mixture for every 100 lb. of dry leather. After fat liquoring, the goods should be horsed up for several hours prior to striking out and drying. The drying should be carried out in a moderately warm, but not hot shed, and it is not advisable to have the goods strained, as it is likely to render the leather hard and impoverished.

When dry, the leather should be stored in damp sawdust for 12 hours or until in the right condition for staking. After staking and drying it requires fluffing on an emery wheel and finally dope finishing

in the usual way.

Belt Dressing

Formula No. 1	
Wool Fat	50 g.
Mineral Oil (0.885-90)	20 g.
Paraffin Wax (56-58° C.)	10 g.
Ceresin, Yellow (58-60°)	5 g.
Castor Oil ("Second Press-	o 8.
ing'')	10 o.
Degras	10 g. 5 g.
No. 2	· 8.
Resin	40 o
Train Oil	40 g. 10 g.
Cotton Seed Oil or Sperm	- ° 8.
Oil, Blown	15 g.
Paraffin Scale Wax	±0 g.
(48-52° C.)	15 c
Mineral Oil (sp. gr. 0.905)	15 g. 20 g.
	- S.
No. 3	
a. Wool Fat, Neutral Tallow	30 g.
a. Tallow	20 g.
b. Graphite, Amorphous Castor Oil	10 g.
Castor Oil	10 g.

Melt up the fats a, stir then into the fusion graphite, and castor oil. Press. The product is soft and like a salve.

Shoe Bottom Dressing		
Montan Wax, Bleached	10	oz.
Paraffin (or Scales), White		
(50-52° C.)	10	oz.
Anilin Dyestuff (Oil Soluble)	2	oz.
Turpentine Oil (or Sub-		
stitute)	54	07.

Patent Leather Dressing Black

Formula	No.	7
TOTHUM	T 4 O.	-1

Celluloid	20	lb.
Castor Oil	5	lb.
Lampblack	5	lh.
Alcohol	30	lb.
Benzine	35	lb.

	No. 2	
Celluloid	110. 2	25 lb.
Lampblack		8 lb.
Nigrosine		1 lb.
Castor Oil		6 lb.
Alcohol		20 lb.
Benzine		45 lb.
20111110	No. 3	20 201
Celluloid		25 lb.
Lampblack		8 lb.
Nigrosine		1 lb.
Castor Oil		8 lb.
Alcohol		25 lb.
Benzine		40 lb.
	Red	
Celluloid		30 lb.
Ochre		5 lb.
Castor Oil		5 lb.
Zinc White		3 lb.
Nigrosine		2 lb.
Alcohol		20 lb.
Benzine		30 lb.
	Blue	
Celluloid		30 lb.
Zinc White		5 lb.
Paris Blue		2 lb.
Castor Oil		8 lb.
Alcohol		25 lb.
Benzine		25 lb.
~	Green	00.33
Celluloid		30 lb.
Zinc White		5 lb.
Schweinfurth	Green	2 lb.
Castor Oil		8 lb.
Alcohol Benzine		25 lb. 25 lb.
Denzine		20 1D.

White Shoe Bottom Finish Gum Tragacanth oz. Water 11/4 gal. Soak and stir until smooth, then add Precipitated Calcium Carlb. bonate 1/4 lb. Titanium Dioxide Oxalic Acid lb. Copper Sulphate lb. Magnesium Sulphate Sal Soda 1b. oz. Water gal.

Black Dye for Leather

The following dye solution is used for the dyeing of the uppers of leather shoes. It will render same black in one application regardless of the previous color.

•		e on one provider c			
	Black Dye	(Alcohol Soluble)	4	oz.	
	Methanol		66	oz.	
	Benzol	and the second	20	oz.	
	Nitrobenzol		10	oz.	

The black dye should be of the acid type such as Calco Condensation Black No. 1601. The solvents are mixed and the dyestuff placed in a cloth sack and suspended in the solvent mixture which is occasionally agitated.

Shoe Luster (Finish)

Water	850	cc.
Ammonia (0.910)	20	cc.
Shellac, Bleached, Finely		
Powdered	150	g.

Let stand cold for some hours; heat the jelly formed to liquefy it.

High Luster Finish

(Water	100 cc.
a. { Borax	25 g.
a. \begin{cases} \text{Water} \\ \text{Borax} \\ \text{Shellac, Bleached} \end{cases}	150 g.
b. Water	700 cc.
c. Turkey Red Oil	50 cc.

Dissolve a, warming up gently without boiling; thin with b, and add c.

Dark High Luster Finish

9	
Ruby Shellac, Powder	150 g.
Water, Cold	850 cc.
Ammonia (0.910)	20 cc.

Soak for 6-8 hours (covered), warm to complete solution (if necessary, add more ammonia). Optional: add dyestuff.

High Luster Finish

Ruby Shellac	150	g.
a. { Water	200	čc.
Ammonia (0.910)	30	cc.
b. Water	550	cc.

Make up a, thin with b.

Liquid Burnishing Wax for	Shoe	Soles
Carnauba Wax	20	oz.
Turpentine	20	oz.
Black Dye (Oil Soluble)	3	oz.
Duponol W.E. or Lohrinol	5	oz.
Ferric Acetate	6	oz.
Glacial Acetic Acid	0.2	oz.
Water	45.8	oz.

Reduce the ferric acetate to a powder and dissolve same in the acetic acid and water mixture. Dissolve the Duponol W. E. in the above solution and heat to about 170° F. Melt the carnauba wax and pour into the turpentine which has been previously heated to about 180° F., dissolve the black dye in this mixture, and then add this latter solution to the former while agitating vigorously. Allow to cool with continued agitation. Du-

ponol W. E. is one of a series of soaps	or
emulsifying agents of the higher alcol	
sulphates which are effective as such	in
an acid solution.	

Preserving Hides and Skins German Patent 617,166

Salt		,	99	
Sodium	Perborate		1	lb.

Conservation of Shoe Soles

Conservation	OT	pmoe	pores	
Melt up:				
Linseed Oil			50 - 60	
Paraffin			40-50	g.
Heat 80° C.				

Treat soles with this mixture after thorough cleaning, 2 or 3 times in 4-6 weeks.

Hardener for Shoe Soles

Rosin, Pale	4 g.
Linseed Oil Varnish	5 g.
Dissolve hot and add:	
Benzoline or Turpentine or	
Mixture	9 g.

Impregnation of Shoe Soles French Patent 750,728

$a. \left\{ egin{aligned} ext{Benzoic Acid} \ ext{Acetone} \ ext{Alcohol} \end{aligned} ight.$	3 g.
a. Acetone	40 cc.
Alcohol	10 cc.
$b. \left\{ egin{array}{l} ext{Oxalic Acid} \\ ext{Aluminum Sulphate} \end{array} ight.$	3 g.
b. Aluminum Sulphate	5 g.
Water	50 cc.

Dissolve a and b separately, mix, add 15 g. of dye to 1 liter; brush on roughened soles.

Preservation and Hardening of Sole Leather

	ed Oil				cc.
Wate	er Glass	(40-45°	Bé.)	4	cc.
Mix brush.	until	emulsifie	d.	Apply	with

Waterproofing Leather Formula No. 1

TOTHICLE TION T	
Gutta-Percha Rape Seed Oil, Boiled Yellow Wax Pig Fat	2 g. 8 g. 6 g. 25 g.
Venetian Turpentine	60 g. 1 g.
Spermaceti	I g.
No. 2	-
Linseed Oil	100 g.
Gutta-Percha	10 g.

a little

Copal Varnish

No. 3		
Amber	380	g.
Linseed Oil, Boiled	250	
Sandarac	30	g.
Turpentine, Venice	60	g.
Turpentine	200	
Tallow	600	g.
Caoutchouc	75	g.
Linseed Oil	300	
No. 4		_
For Hunting Shoes		
Caoutchouc	4	g.
Pig Fat	6	g.
Cod Liver Oil	24	g. g. g.
No. 5		
For Horse Covers		
Japanese Train Oil	94	g.
Saturated Caoutchouc Solu-		-
tion in Turpentine	5	g.
Aniline	1.5	g.

Quick Black Shoe Edge	Ink
Bright Drving Carnauba	•
Bright Drying Carnauba Wax Emulsion	50 lb.
Nigrosine	8 lb.
Water	3 gal.

Edge Filler for Shoe	Factory Use
Soap	15 lb.
Yellow Dextrin	5½ lb.
Neatsfoot Oil	1½ qt.
Oil of Mirbane	1 pt.
Gelatin	11½ lb.
Formaldehyde	1 qt.
Water	1 at.

This is made up with sufficient water to make 60 gal. solution.

Brown Shoe Heel Stain

Alcohol	7 fl. oz.
Acetone	1 fl. oz.
Gum Tragacanth	4 oz.

Mix the above until gum is thoroughly wetted and to it add slowly with stirring the following solution made by boiling and then cooling:

Oxalic	Acid			3	oz.
Water	Soluble	Brown	Dye	8	oz.
Water				$2\frac{1}{4}$	gal.
		_			-

Strain through cheesecloth.

Shoe Dye Remover

Isopropyl Alcohol	. 7	CC.	
Acetone	1	cc.	
Butyl Cellosolve	1	cc.	
Water	10	cc.	

Shoe Repairing Cement U. S. Patent 2,004,059

Six pounds crepe rubber, 2.5 lb. rosin, and 1.5 lb. zinc dimethyl dithio carbamate, said components fluidified in 15 gal. of benzol.

Fat Liquor, Leather

Lecithin	50	lb.
Water	50	lb.
Soda Ash	1/2−1	lb.

Mix the above well and then mix in a suitable quantity of neatsfoot oil.

Russia Leather from Rejected Hides

The washed and pressed leather is greased in a drum with a mixture of 2 kg. train oil, 5 kg. mineral oil and 4 kg. degras per 62-5 sq. m. of hides, drummed 40 minutes while warm, spread, stoned, dried for 4-5 hours to 38-40% water content and cut through the middle into halves. The damaged spots are cut out, the hides reset and greased by hand on both sides with a mixture of degras 2 kg., train oil 6 kg., mineral oil 6 kg., lard 6 kg. and tar 5 kg. per 100 sq. m. The leather is left for 12 hours and dried at 28-30° C. to a water content of 32-5%. left for 6 hours to assure a uniform distribution of the water and finally worked over with the whitening sleeker. leather is then dyed, greased on both sides with a mixture of 3 kg. train oil, 4 kg. tar, 6 kg. mineral oil and 2 kg. paraffin, allowed to rest 12 hours, dried at 28-30° C. and treated with a mixture of 150 g. nigrosine, 125 g. gum tragacanth, 50 g. carpenter's glue, 1.5 liter blood and 1 liter milk (all mixed with 12 liters water). The goods are finally dried, polished and sorted.

Preserving Lizard Skin

Skins are treated with	
Zinc Chloride	2 lb.
Salt	10-20 lb.
Water	100 lb.

Protection of Hides and Skins from Skin Beetle

Salt thoroughly applied to hides gives excellent protection against beetle attack. Heavily salted hides which are first rubbed with salt and then soaked in saturated brine for 10 hours or are merely soaked in the brine, are entirely protected during storage for 6 months in the summer in a beetle-infested room. Hides which are rubbed on the flesh side are not so well protected. Hides are protected almost completely by dipping

them, immediately after flaying, in a 2.5% sodium arsenite solution. Spraying sun dried hides on the inside with the sodium arsenite solution does not altogether protect the grain, although it does so to some extent. Sodium arsenite has a marked preservative action on the hides, but a solution stronger than 2.5% is required to prevent decay when hides are dried in the shade in humid regions. When they are stored with salted and untreated hides, the sodium arsenite treated hides do not act as a bait for the beetles and no dead insects are found on them. The sodium arsenite treatment has no deleterious effect on the leather prepared from the hides, and the workmen who handle the hides show no signs of arsenical poisoning.

Stuffing for Welting Leather
Cod Oil 1 gal.
Sulphonated Cod Oil 1 gal.

The above mixture is used per 100 lb. of welting.

Tanning Shearlings

Soaking: Skins are soaked in clean water, salted skins 10 to 24 hours; dry skins several days, according to condition. Skins must be thoroughly soaked but care must be taken that the wool does not become loose. To prevent this different ingredients are added to the soaks. Small quantities of any of the following may be used: zinc chloride; formaldehyde or alum.

Naphtha or degrading compounds are the most efficient for removing the excess grease; these being reclaimed by distillation and the grease is recovered as a byproduct. In case the stock is not degraded it should be thoroughly washed with a warm soap and soda solution. After degreasing all burrs and brands are worked out. Neglecting to clean out burrs will cause damage in the unhairing machine. Skins are then washed by hand to remove all dirt and to render them as white as possible. This step in the process may be accomplished in the paddle or drum which has a tendency to loosen the wool.

The pickling or tanning may be carried on in the paddle or by hand. If the paddle is used a base solution of approximately 1 lb. of salt for each gallon of water is used and then built up to the desired salometer with equal parts of salt and alum. This amount should be about 4% of each on the weight of stock. This solution may be used several times by the addition of equal parts of salt and alum figured on the weight of the stock.

A small amount of sulphuric acid may be used if desired. This bath is worked up to 50 or 60° C. over a period of 3 days. When stock is struck through it is taken out and drained and is ready for oiling or may be retanned with gambier or quebracho. White and light shade stock is finished out of the alum.

Skins tanned by hand are best treated on the flesh with salt and sulphuric acid solution. This solution is made with 1 lb. of salt and 1½ oz. of acid to each gallon of water. This solution is applied to the flesh with a brush and the skins piled flesh to flesh or folded down the back with the flesh side in. The next morning the stock is given an alum tan on the flesh made up as follows:

Alum	5%
Salt	5%
Sodium Bicarbonate	5%
Flour	5%
Egg Yolk	1%
Oil	1%

The flour should be worked into a paste, after which the other ingredients are added, the egg yolk being dissolved in a small quantity of cold water. Soda should be added slowly. Two coats of this mixture are given at intervals of 10 to 12 hours at which time stock should be thoroughly tanned. Stock is now thoroughly dried out after which it is sammied back, staked and a light coat of oil given the flesh or a fat liquor may be given, made up of soap, neatsfoot oil and sulphonated oil. Stock before becoming thoroughly dried is staked and stretched. After skins are dried they are restaked, snuffed, combed and clipped. If desired stock can then be dyed or bleached.

Russia Leather Odor
Bases for this odor are:
2-Tertbutyl 4,5 dimethyl-1-phenol.
or
2-Isopropyl-4,5-dimethyl-1-phenol.

LUBRICANTS, OILS, FATS

Gear Lubricant for Arctic Climates

In the northwestern section of the United States and a large section of Canada air temperatures of 40° below zero are not uncommon. At temperatures such as these ordinary winter gear oils are too viscous to permit satisfactory operation of motor cars, and many motor car manufacturers have recommended diluting the gear oil with kerosene to meet these conditions. This practice has always been frowned upon by lubrication engineers since even if the lubricating value of the oil is not entirely destroyed by such dilution, the facilities of the average service station for accurately blending without danger of contamination are not the best. The following formula will produce a lubricant which will give satisfactory performance and adequate lubrication under arctic weather conditions.

Thickened Rape Oil	8 lb.
Asphaltic Black Oil	7 lb.
(90 visc. at 210° F.) Gulf Coast Pale Oil	85 lb.
(100 visc. at 100° F.)	

Sulphur Lubricant Base

The use of sulphur for manufacturing lubricants of high film strength is rapidly gaining popularity. The formula given here will produce a base which can be diluted with mineral oils to make cutting oil and various extreme pressure compounds.

Flowers of Sulphur	10 lb.
Lard Oil	90 lb.

Mix well and slowly raise the temperature to 425° F. Maintain mild agitation throughout.

Anti-Rust Compound

Rust and corrosion will do more damage to machinery than several months of hard service. This is particularly true of construction and railway machines which must often be left exposed for long periods. A simple formula for an efficient and economical protective compound is given. The materials should be heated, mixed well and applied with an old paint brush.

Paraffin Wax	6	lb.
Asphaltic Still Residue	94	lb.
(About 1000 visc. at 210	° F.)	

Steering Gear Lubricant

With the general trend to wider treads on automobile tires it has been necessary to redesign steering gear mechanisms to avoid hard steering. Automobile engineers agree that special lubricants are required for most efficient operation.

Oleic Acid	300	lb.
Lime	43	lb.
Water	16	gal.
Western Cylinder Oil	475	
Sulphur Base	1000	ĺb.

Proceed the same as for making lime soap grease except that the sulphur base is not added until the other ingredients are completely cooked.

Mixed Base Grease

The following formula will make a grease which combines the advantages of the smooth texture of calcium soap grease with the cohesive rubber-like character of aluminum oleate. Although the melting point of this grease is not materially higher than a similar calcium soap grease, the melted grease has the slow flowing characteristics of aluminum greases. The formula given is for a medium consistency but other grades can be made by varying the soap content.

Lime	17 lb.
Fat	113 lb.
Aluminum Oleate (Pulp	
Stock)	50 lb.
Pale Oil (100 Viscosity)	112 gal.
Water	6 gal.

Place the fat in a steam jacketed kettle equipped with paddles for stirring, add a small portion of the mineral oil, mix the lime with sufficient water to form a thin paste and add this to the material in the kettle. Turn on the steam and start the paddles. When the soap has cooked for 5 hours it should be tested to determine if saponification is completed, if so the steam is turned off and half of the balance of the mineral oil is run in slowly. The rest of the mineral oil is run into a separate kettle and the aluminum oleate melted in it and this mixture is pumped into the first kettle while still warm. Stirring should be continued until a smooth uniform grease is produced.

Non-Bleeding Grease

One of the difficulties encountered in the use of pressure grease is the tendency of the light oil to separate and bleed away leaving the bearing choked with a hard soap. This formula produces a grease which will stand indefinitely without separating. This is not a high melting point grease and is intended for automobile chassis lubrication and similar applications.

Green Petrolatum	250 lb.
Paraffin Pale Oil (28° Bé.)	92 gal.
Lime	9 lb.
Fat	55 lb.
Water	3 gal.

Melt the petrolatum in the mineral oil. Mix well, then proceed as for ordinary calcium soap grease.

Lubricant for Bearings with High Temperatures and Pressure

Formula No. 1 Rosin Wool Fat Stearin Mineral Oil (0.900-7) 80 g. Castile Soap 15 g. Caustic Soda (40° Bé.) 4 g.

No. 2		
Rosin	5.5	g.
Wool Fat, Crude	6	ğ.
Wool Fat, Stearin	11	ğ.
Tallow	5	ğ.
Linseed Oil	5	g.
Caustic Soda (35° Bé.)	5	g.
Mineral Oil (0.885-90)	78	ğ.

No. 1 is a high melting fat (150-200° C.), No. 2 melts at about 100° C.

The saponification is done in a directly heated kettle (cast iron), which has a removable stirrer, at 150-200° C. Test: should not sweat oil or alkali when pressed with the finger after cooling. If desired, short-cut fibers may be added to the mass. Solidify in patterns and cut into briquets.

Metal Rolling Lubricant

Tallow	60	lb.
Yellow Soap	15	lb.
Water	92	gal.
Heat and stir until smooth.		

Non-Greasy Lubricant U. S. Patent 1,970,902

Sodium	Alginate	19	oz.
Water		100	oz.

Mix to a smooth paste while heating to 100° C. Add

Glycerin 76 oz.

Boil off nearly all of the water.

Olive Oil Motor Lubricant

Olive Oil (Low Titre) Mineral Oil 75 fl. oz.

Lubricating Grease for Carriages

Blue Oil	45 g.
Slaked Lime	6 g.
Rosin Oil	22.5 g.
Fat Soluble Black Dye	0.2 g.
Dissolve in the blue oil	

Chain Lubricant Formula No. 1

Stearin	85 g.
Beeswax	5 g.
Carnauba Wax	5 g.
Japan Wax	5 g.
No. 2	· ·
Stearin	85 g.
Beeswax	2.5 g.
Carnauba Wax	5 g.
Janan Wax	5 0

Pour at lowest possible temperature and allow to cool slowly and undisturbed.

Penetrating Oil British Patent 414,847

Useful	for	loosening	rusted	meta.
parts.		_		
Engine	Oil		1	qt.
Naphth	a or	Kerosene	3	qt.
Carbon	Disu	lphide	2	oz.
Oil of (Campl	nor	1-2	oz.
Graphit	e. Po	wder	1-4	oz.

Core Oil

Formula No. 1		
Linseed Oil	300	oz.
American Gas Oil	600	oz.
Dark Whale Train Oil	100	oz.
No. 2		
Rosin	200	oz.
Train Oil	200	oz.
Vulcan Oil	600	OZ.

Stuffing Grease

	(Melting Po	int	over	96°	C.)	
. !	Tallow			. 3	2	g.
a	Tallow Lard Oil				3	g.
ъ.	Lime Hydrat	e			2.5	g.
c.	Zinc Oxide				2.5	g.

d. Machine Oil, Refined 4-5° E. Viscosity 50° C. 79 g.

e. Yellow Aniline Dye, Oil
Soluble
f. Water

0.03 g.
1 g.

Notes: Lime Hydrate—Made up of finest commercial lime hydrate, diluting with water 1:4.

Work bringing a into kettle with 1/3 to 1/2 of needed d; heat to 80-90° C., add slowly b, continue warming. At 100° C the mass starts "rising" in the kettle, then diminishes when water is evaporating.

Tests: Should be resistant against not too strong finger-pressure; weakly brittle, should not sweat out water or oil when cooled. On the other hand, a water insufficiency is indicated if mass is too brittle (in this case add little boiling water). If tests are satisfactory, add the remainder of d, at 70° C. or warmer—not too slowly, not too quickly. The aniline dye dissolve in mineral oil.

Let stand over night. Stir till cool next day.

Cutting Oil Formula No. 1

a. Mineral Oil (Spindle Oil)80 g."Tall-Oil," Refined20 g.b. Caustic Potash (40° Bé.)6 g.c. Methylhexalin1-2 g.Saponify a with b, clear with c.No. 2

 Paraffin Oil (28 to 30° Bé.)
 250 g.

 Rosin
 22 g.

 Oleic Acid
 22 g.

 Caustic Soda
 3 g.

 Water
 10 g.

 Alcohol
 7 g.

No. 3
Lard Oil (No. 1) 1 gal.
Paraffin Oil (28° Bé.) 52 gal.

Manipulation: Mix at room temperature.

No. 4

Lard Oil (No. 1) 5 gal. Extra Lard Oil 5 gal. Paraffin Oil (28° Bé.) 42 gal.

Manipulation: Mix at room tempera-

Non-Corrosive Cutting Oil U. S. Patent 1,979,250

	- 000110		
Mineral Oil		71-74	lb.
Castor Oil		814-91/2	lb.
Rapeseed Oi	1	81/4-91/2	lb.
Caustic Pot	ash	814-91/2	lb.
Soda Ash		0.6-11/4	
Mix and dil	ate with	water.	

Brake Oil (Non-Rancid)

a. Mineral Oil (Spindle Oil) 1000 g.b. Paratoluol Sulphochloride 5-6 g.

a. Rape Seed Oil 900 g. Camphor Oil 100 g. b. Paratoluol Sulphochloride 5-6 g.

Dissolve b in little part of a, then add to the above amount.

Gasoline Motor Lubricant British Patent 423,441

Mineral Oil 99 lb. Chromium Oleate 1 lb.

Radiator Anti-Rust Compound

In the past year the automotive industry has given much attention to the prevention of rust and corrosion in automobile cooling systems. Engines with aluminum composition cylinder heads have received the most attention but even in the case of ordinary steel parts it has been found that cooling systems are more efficient if rust and scale formation is prevented.

For this purpose soluble cutting oil such as is used for machining metal is very efficient. The only limiting factors are acidity and alkalinity. Soluble oils having a high acidity will corrode the radiator while too much free alkali will damage aluminum cylinder heads. Several of the formulæ given in volumes one and two of THE CHEMICAL FORM-ULARY will be very satisfactory as cooling system corrosion preventatives. The usual quantity used is ½ oz. of soluble oil for each gallon of water.

Greaseless Lubricating Pencil

Useful for lubricating hinges of automobile doors, etc., as it will not run off and produce stains or accumulate dust.

Beeswax 80 g.
Diglycol Stearate 20 g.
Graphite Powder 100–200 g.

Melt together and stir until just cold enough to pour. Pour into molds and allow to set.

Dynamo Brush Lubricant

Ceresin	20 g.
Tallow, Acid Free	10 g.
Wool Fat, Neutral	10 g.
Castor Oil	10 g.
Vaseline Oil	50 g

Melt together and add enough organic solvent (Heavy Benzoline, Naphtha or Tetralin).

Cotton Spindle Machine	Oil
Spindle Oil, Refined (5-6°	
E at 20° (C.)	85 gal.
Rape Seed Oil	15 gal.

Veneer Press Caul Lubricant German Patent 596,345

Neutral Soap	15	oz.
Lanolin	35	oz.
Petrolatum, Liquid	30	oz.
Formaldehyde	16	oz.

Transformer Oil U. S. Patent 1,988,299

Crude Mineral Oil 99.5 g. Phenyl Alpha Naphthylamine 0.5 g.

Transformer Oil Canadian Patent 353,332

To a mineral oil of iodine value of 7 to 20 about 0.5% phenyl α -naphthylamine is added to retard sludge formation.

Petroleum Proof Valve Lubricant Citric Acid, Anhydrous 64 g. Tetraethylene Glycol 97 g.

Heat at 180-185° C. for 90 minutes; cool. Do not overheat or an infusible product will form.

Rubber Mold Lubricant

Cocoa soapstock, a material containing a large percentage of coconut oil saponified with alkalies to give a pure hard soap, makes a suitable product for lubricating molds to prevent sticking of the vulcanized stock. If properly made, without traces of sodium silicate, it will not cause caking on the molds. The recommended quantity is 8 to 12 lb. to a 55 gal. drum of water. The soap is dissolved in water by cooking, either by open steam or external heat of some kind. For easy spraying the solution is kept warm by steam or a small electric heating unit can be applied at the spray nozzle to prevent clogging.

Screw Thread Lubricant

Flaked graphite mixed with a medium grade of lubricating oil to form a paste and applied to the threads of screws and bolts facilitates the backing off of nuts and the removal of screws and machine bolts. The paste, which also is suitable for pipe joints, prevents rust.

Vacuum Tap Grease

Rubber	30	oz.
Rosin	15	oz.
Pine Pitch	50	oz.
Soot	5	oz.

Journal Grease U. S. Patent 1,989,196

Heavy Black Petroleum Oil	5.9 lb.
Heavy Steam Refined Pe-	
troleum Oil	34.4 lb.
Stearic Acid	40.5 lb.
Caustic Soda (48° Bé.)	13.1 lb.
Lard Oil	6.1 lb.

Spring-Leaf Lubricant British Patent 414,948

White Lead in Linseed	Oil	
(92 Lead, 8 Oil)	83-84	lb.
Graphite Powder	5.2	lb.
Petroleum Grease	10.4-10.5	lb.
Glycerin	0-1.3	lb.

Nickel and Monel Drawing Lubricant

A paste made of castor oil and lead, recommended for use as a lubricant in the cold forming of Monel metal and nickel, can be removed by a number of solvents. Carbon tetrachloride, being non-inflammable, is to be preferred. Benzene, gasoline, and alcohol produce satisfactory results

factory results.

Cold soap and caustic solutions are not entirely satisfactory but can be used as an alternative, if necessary, when they are used hot.

Wire Drawing Lubricant U. S. Patent 1,944,273

Sodium Alginate	1	lb.
Tallow	4	lb.
Soap	2	lb.
Water	195	lb.

Drawing Die Lubricant for Diamond Dies

Rye Flour	6	lb.
Water 10	00	lb.
Beef Tallow	$2\frac{1}{2}$	lb.
Soft Soap	21/2	lb.
Heat and stir until uniform		

Corrosion Protecting Grease

Neutral I	Petroleum	Grease	100	oz.
Zinc Chro	mate Pow	der	$-2\frac{1}{2}$	oz.
Pyridin I	Bases (Cr	ruđe)	1	oz.
Durb tomat	than to fo		th amo	

Rub together to form smooth grease.

Lubricant for Preventing Corrosion French Patent 778,792

Sodium Peroxide	1/4	oz.
Methanol	2	oz.
Hydrogenated Phenol	4	oz.
Lubricating Oil	100	OZ.

Lubricating Haulage Ropes

Before the lubricant is applied, the surface of the rope should be cleaned and dried, because oil or grease applied to the surface of a rope covered with mud or coal dust, water and old oil will be thrown off without having the slightest chance of penetrating to the interior. In most cases the treatment can be given to the rope during an idle shift.

Main ropes used on inclines can be treated as follows: The rope should be wound very slowly on to the drum, the surface being cleaned as it enters the engine house. Cleaning should be done with wire brushes without using a solvent, such as petrol or paraffin. brushes may from time to time be washed in paraffin, but this should be shaken off before using the brush on the rope again. The cleaning may be completed with waste or sacking. No solvent (petrol or paraffin) should be used on the rope, because experience has shown that the solvent readily penetrates into the middle of the rope and rapidly dissolves out any remaining lubricant. The rope should be allowed to remain on the drum long enough to allow it to dry as much as possible.

When the rope has been cleaned and dried, the lubricant should be applied by hand with a fairly stiff brush. Devices in which the rope is caused to pass under a roller in a bath of oil are less effective and are wasteful. It is important that the rope should be dry when the lubricant is applied otherwise the oil will not adhere, and the work should be done within the engine house as the rope leaves the drum. If the lubricant is applied in the open, a shower of rain may render useless the whole operation of cleaning and drying the rope. The successful lubrication of a haulage rope calls for a good deal of skill and patience, but unless it is properly done the time and materials are wasted. It is better to do a portion of the rope well each week than to waste a lot of grease by applying it to the whole of the rope without cleaning and drying.

It is not possible to lay down any fixed periods for the lubrication of haulage ropes, because the periods will vary with the working conditions. A rope oil.

which makes a large number of journeys on a wet incline will need lubrication every week, whereas a rope which makes only a few journeys in the dry may be kept in good condition by less frequent treatment. Excellent results have been obtained on endless rope haulages where the rope is lubricated continuously. In one instance a light mineral oil is allowed to drip on to the moving rope at the rate of one drop per yard; this rope works on a comparatively clean and dry roadway.

Research is in progress as to the best type of oil for applying to ropes in service. At the moment it would seem that the best results are obtained with a medium heavy mineral oil. The oil must be free from acidity, and should contain

no filler or soapy material.

Hot Neck Grease

Asphaltic Residue	10 lb.
Candle Tar Pitch	20 lb.
Paraffin Cylinder Stock	
(700 Fire Test)	70 lb.

Heat to 550° F. and blow with air until melting point of 200° F. is obtained.

Above is cast into blocks and used for the lubrication of roller necks in steel mills.

High Temperature Lubricants British Patent 431,066

Lubricants for use at high temperatures, e.g., in internal-combustion engines, consist of lubricating oil in which is dissolved or dispersed chromium or an organic compound thereof, and one or more other substances preventing sludging, e.g., organic compounds of tin and/or lead. Up to 1% of each addition is suitable. For example, 0.5 lb. of chromium oleate, 0.1 lb. of tin oleate, and 0.1 lb. of tetraethyl lead are added to 100 lb. of a compounded vegetable and mineral lubricating oil; or 0.4 lb. chromium oleate and 0.1 lb. of tin oleate to 100 lb. or a paraffinic mineral oil.

Non-Chilling Lubricants Formula No. 1

Mix
Castor Oil 3 cc.
Paraffin, Chlorinated (30%
Chlorine) 7 cc.

Spindle Oil, Russian 190 cc.
This gives a highly cold-resistant, clear

No. 2		Turpentine Ammonia (28%)	8.7 lb. 4.4 lb.
Castor Oil Paraffin, Chlorinated (30%	10 cc.	Graphite Powder	30 lb.
Chlorine) (Heat to 200° C.)	10 cc.	Watersoluble O	il
Spindle Oil, Russian	80 cc.	Naphthenesulphonic Acid Olein (or Liquid Wool	s 15 g.
No. 3		Fatty Acid)	5-7 g.
Spindle Oil, Russian Paraffin, Chlorinated (40%	40 cc.	Spindle Oil, Refined (60° Caustic Potash (25° Bé.)	until neutral
Chlorine) Castor Oil	40 cc. 20 cc.	Hexalin and Tetralin (1:	1) 3–4 g.
***************************************	40 CC.	Mineral Oil Soluble Ca	stor Oil
Rod Lubricant a. Ceresin, Yellow	95 ~	To obtain castor oil w	
Sperm Oil	25 g. 25 g.	soluble in mineral oil, heat the former with 30 parts	of trichloro-
Tallow Melt together.	50 g.	ethylene for 2 hours in a cl 130° C. The pressure will	
or		atmospheres. After distilli solvent, the resulting casto	
b. Ceresin, Yellow Spindle Oil, Refined	1 g. 3–8 g.	soluble in mineral oil. The not be brought about by he	is result can
Melt at low temperature.		alone or by refluxing with	solvent. A
Solid Lubricant		second method is to heat in 90 parts of castor oil with	. 10 parts of
Formula No. 1		carbon tetrachloride for 2 h The pressure increases to	
Canadian Patent 344, Heavy Distilled Naphthenic	966	atmospheres. Dissolve in m distil off excess solvent, rem	ineral oil and
Petroleum	30.8 1ъ.	traces by distillation in vac	10.
Residual Naphthenic Petroleum	13.6 lb.	Tuhmisant Tussluble in Our	onio Solmanto
Stearic Acid Oleostearin	14 lb. 28 lb.	Lubricant Insoluble in Org Mix to a paste the follow	
Caustic Soda Water	6.6 lb. 7 lb.	Anhydrous Glycerin Dextrin	25 oz. 7 oz.
No. 2		Pure d-Mannitol	3.5 oz.
Canadian Patent 344,	967	Heat carefully with consuntil the solid material is	stant stirring
Viscous Naphthenic Petroleum	43 lb.	the solution begins to boil,	then cool to
Animal Fat Aluminum Stearate	39.4 lb. 4.7 lb.	room temperature with stir crease the viscosity, add n	nore dextrin;
Caustic Soda Slaked Lime	5.3 lb. 0.6 lb.	to increase fluidity add more increase greasiness add mor	
Water	7 lb.		
Hard Grease		Tempering Fats (Bath to Harden Steels)	
Train Oil Fatty Acid Lime, Hydrated	12 g. 2 g.	Formula No. 1	
Zinc Oxide Spindle Oil	2 g. 82 g.	Peruvian Bark Powder	500 g.
Water	2 g.	Neatsfoot Meal Salt	500 g. 850 g.
Melting point 75° C.		Saltpeter Potassium Ferrocyanide	250 g. 15 g.
Graphite Lubricant	8	Soft Soap	1000 g.
U. S. Patent 2,003,56	34	No. 2 Beef Tallow	10 g.
Degras (Free from Fatty Acids)	20 lb.	Potassium Ferrocyanide, Powder	
Kerosene Water	16 lb. 75 lb.	Wax Colophony (Rosin)	2 g. 2 g. 2 g.
	.0 10.	Combiner (100sin)	4 g.

Waterproofing, Perilla Oil

One method is to react one part of straight phenolic resin with 2 or 3 parts of perilla oil at between 500° and 550° F. If polymerized perilla oil is used, even better results are obtained. Another method is to employ some wood oil. For instance, one part of straight phenolic resin to 2 parts of wood oil may be reacted together and then extended with various amounts of polymerized perilla oil. Another formula is phenolic resin 5 parts, wood oil 10 parts, perilla oil 85 parts. Another is 10 phenolic resin, 20 wood oil, and 70 perilla oil. All parts are by weight.

Coloring Lubricating Oils British Patent 424,205

Lubricating oils are improved in color by adding a solution in mineral oil or other blending agent of the product obtained by heating together until fluorescence develops, an aeridine, rhodamine, eosine, or eurhodine dye with stearic acid and a water-insoluble soap. Soaps specified are aluminium stearate, magnesium stearate, oleate, or resinate, and zinc soaps. For example, 1 lb. of phosphine 5G., 1 lb. of stearic acid, and 3 lb. of aluminium stearate are heated to 120° C. until the fluorescence is a maximum; the mixture is cooled, pulverized, and dissolved to a 10% solution in a mineral oil miscible with lubricating oil. 0.25–0.5 gal. of the solution is added to 100 gal. of lubricating oil.

Refining Lubricating Oil U. S. Patent 2,020,954

Stock of about 68 viscosity index is subjected to the simultaneous action of 10% of aluminum chloride and 10% of fuller's earth at a temperature of about 350° F. for ½ hour.

Purification of Lubricating Oil

If lubricating oil is shaken with phenol, the lower layer consists of oil and impurities in phenol; the upper layer consists of phenol dissolved in pure oil. The phenol is removed and recovered by distillation or by washing with sulphuric acid.

Dewaxing Mineral Lubricating Oil U. S. Patent 2,014,629

Amorphous wax is eliminated by treating the wax bearing oil with 3 to 10% of substantially anhydrous aluminum

chloride at a temperature of about 200° for a half to four hours, thinning with a light distillate, chilling and filtering.

Dewaxing Oil U. S. Patent 1,978,010

A process for treating wax-oil mixtures comprises mixing 1 to 4 volumes of methylene chloride with 1 volume of the mixture, chilling the mixture to a temperature below 0° F. and filtering precipitated wax from the mixture.

Preventing Discoloration of Oils and Fats

British Patent 410,834

Discoloration of animal or vegetable oils or fats on exposure to air and light is prevented by incorporating not more than 0.05% of colloidal copper, cobalt, cadmium or silver, or of the carbonate of cobalt, copper, lithium, manganese, cadmium, barium, bismuth, the nitrate of calcium, beryllium, or lithium, the acetate of sodium, copper, manganese, the hydroxide or cobalt, beryllium, copper, thorium or of a mixture of cobalt carbonate and copper carbonate with or without bismuth subcarbonate.

Reclaiming Used Lubricating Oil U. S. Patent 1,936,901

Used Lubricating Oil	100	gal.
Red Oil	1	gal.
Calcium Hypochlorite	6-8	gal.
Sulphuric Acid	6	lb.
Mix together and then add:		

Sodium Silicate 50–100 lb. Water 10–20 gal.

Heat at 52-122° C. for two hours. Cool; add water, 3 gal., and separate clear oil.

Fat and Oil Bleaching

In refining fats and oils the color is improved by adding 8 to 10% soap stock to the fat.

Decolorizing Tea Seed Oil

Kaolin 25 lb. Animal Charcoal 20 lb.

The above mixture has been found to give the most economical results.

Increasing Viscosity of Mineral Oils British Patent 416,513

Thickened mineral oils which form gels at room temperatures are obtained by

dissolving less than 2% of cellulose stearate or palmitate in the heated oil.

Oil Filter Mass U. S. Patent 1,940,317

Cotton Waste Curled Hair

75 oz. 25 oz.

Fat Hydrogenation Catalyst
The catalyst is prepared as follows:
Precipitate a solution of 160-300 g. per liter nickel sulphate with a 15° Bé.

sodium carbonate solution at not over 32-65° C., filter on a filter press, wash till free from sulphates with water at 30-50°, dry 4 to 5 hours at 100-105°, grind, sieve, mix with sunflower seed oil and reduce by heating the oil in presence of hydrogen; time of reduction is 5 hours; the temperature is raised to 170-200° during the first hour, to 200-240° during the next two hours and to 240-245° during the last 2 hours. Reduction of the catalyst can be carried out in the same autoclave as the subsequent hydrogenation. The activity of the catalyst lasts over a prolonged period.

MATERIALS OF CONSTRUCTION

Metal Cleaning

Many "mysterious" finishing troubles are due to improper cleaning. What cleaning materials and methods to select will depend upon: (1) the size and character of articles to be cleaned, (2) their surface condition, (3) the volume of work to be handled, (4) the kind of finish to be applied, and (5) various conditions peculiar to the particular factory department wherein the cleaning is to be performed.

Rust, dust, greases, and grit can be cleaned off metal surfaces by the use of one or more of several methods. They may be burned off, chemically removed with an acid or an alkali solution, absorbed by gasoline or naphtha, buffed, or

removed by sandblasting.

Old varnish or paint may be removed by the burn-off process, preparatory to refinishing. A temperature of 650° to 700° F. is required to dislodge the old coating which can then be wiped off with a rag while still hot. The burn-off (oven) process is also a means of drying washed and chemically treated parts.

Heavy rust spots are usually removed by wirebrushing, sandpapering or sandblasting. Thin coatings of rust may be removed either by kerosene or gasoline or by pickling in a solution made of commercial sulphuric acid diluted in water. Other solutions used are: (1) A 20% solution of sodium citrate and water, (2) a 10% solution of ferrous sulphate and water, and (3) a 3½% solution of boric acid and water.

Aluminum parts are prepared for a baked finish by a thorough cleaning with gasoline or naphtha, and a subsequent oven-drying. Old paint and varnish may be removed from aluminum with any standard paint or varnish remover.

Metal Cleaning Composition Canadian Patent 345,172

A compound containing trisodium phosphate and sodium dichromate is used for cleaning tin-coated metal. It inhibits checking or spangling. A satisfactory composition contains trisodium phos-

phate 55 lb., sodium carbonate 40 lb., and sodium dichromate 5 lb.

Cleaning Metal Before Painting

Apply
Ammonia (28%) 1 l.
Alcohol 26 l.
Water 25 l.

Wipe off metal thoroughly after application.

Cleaning Iron and Steel U. S. Patent 1,943,875

Prior to galvanizing or tinning the metal is exposed to the fumes of 1 to 2% of phosgene at $100-200^{\circ}$ C.

Cleaning Tin Surfaces

a. A bath is made up of palm oil that has been heated to 300° F. Any method of heating may be employed as the flash point of the palm oil is quite high. Generally speaking, there is no danger of overheating. Probably the most practical method of heating is by using a steam coil in the palm oil container, as the temperature may be easily controlled.

The work is dipped into the solution of heated palm oil for two to three minutes and removed. No further processing is required for the palm oil is quite liquid at this temperature and will flow freely from the work. It may be found necessary to remove some of the oil by using an air blast to blow the oil from

the work.

The method suggested above will operate well on small work. However, if the work is large, it may be necessary to preheat the work before immersing it into the oil bath. Without preheating heavy work, the oil will cool too quickly when the work is being removed from the solution and will leave an unsatisfactory waxy deposit on the work. The preheating is best accomplished by immersion in superheated water long enough to heat the work sufficiently. Upon removing the work from the heated water, it may be immersed immediately in the palm oil bath.

 Another method that may be used with good results is to immerse the work in a 2% solution of water and nitric acid. This procedure is most efficient if the work is first preheated in water as suggested in Method a. The acid dip is immediately followed by immersion in a rinse of kerosene oil. The duration of the acid dip must be found by experiment as the length of dip depends upon the thickness of the oxide. This may be easily determined by the trial and error method. Too short a dip does not restore the luster, and too long a dip increases the tarnish and produces a spangle effect as in galvanizing. acid dip and kerosene rinse are operated at room temperature.

The drying of the work is best accomplished by drying in heated sawdust. Care must be exerted in this operation as machined work will rust if it is not dried

thoroughly and quickly.

The success of cleaning of tinned work depends upon the quality of the tinning that was on the work originally. It is impossible to produce a luster on an article that had a poor finish in the first place.

Cleaning Monel Screw Machine Parts
The use of sulphur base cutting oil in
high speed automatic screw machine operations, may discolor the Monel metal
parts. This discoloration is due to the
formation of metallic sulphides by the

sulphur in the oil.

The discoloration is readily removed by dipping the parts in a cold solution of sodium cyanide. The solution is made up in the proportions of water 1 gal, sodium cyanide ½ to 1 lb. The time required for cleaning is from 5 to 30 minutes, depending on the degree of discoloration. Caution should be used in handling this solution as it is a deadly poison.

Coloring Metals

Metals are colored chemically or electrochemically by producing thin films of oxide, sulphide, phosphide, silicide, nitride and carbon on their surface. For quantity production, coloring is usually carried on in a rotating drum, while large pieces and objects of art are treated by hand. A few recipes follow:

1. For Copper

a. Brown: immersing in molten sodium nitrate, or imbedding in a paste of 15 parts ammonium carbonate and 5 parts each of copper acetate, tartaric acid in

vinegar, and salt; another solution is 25% copper sulphate, 25% nickel sulphate, 12% potassium chlorate, 7% potassium permanganate.

b. Gray-black: a hot watery solution of 12% copper sulphate and 1% potas-

sium permanganate.

c. Black: 40-50° C. (104-122° F.) warm solution of 600 g. copper nitrate in 200 g. water and 2.5 g. silver nitrate in 10 g. water is brushed on the object and dried at 230° C. (446° F.); or a solution of 10% sodium chlorate, 5% caustic soda and 10% potassium persulphate is used for immersion.

d. Green patina: solution of 25% ammonium chloride, 25% ammonium carbonate, or an acetic acid with an addition

of 1-2% tartaric acid.

e. Blue: 80° C. (176° F.) hot solution of 13% thiosulphate and 3.5% sugar of lead, or of 100 g. potassium chlorate, 100 g. ammonium nitrate and 1 g. copper nitrate in 1 l. water. The objects are immersed for 5-10 minutes.

f. Purple-gray: immersion in a solution of antimony trichloride in water with an addition of equal weight of 5% hy-

drochloric acid.

2. For Zinc

- a. Yellow: aqueous solutions of 5% copper sulphate, 5% sal ammoniae and 3% ammonium chloride are brushed on.
- b. Black: solution of 16% copper sulphate, 8% potassium chlorate in 1 l. water; or a cold solution of 8 parts hydrochloric acid, 3 parts copper chloride, and 2 parts copper nitrate in 64 parts of water.
- c. Iridescent: immersing in a solution of 3 parts tartrate of copper oxide and 4 parts of caustic soda in 48 parts of water. According to duration of immersion, purple, blue, green, yellow or red hues are obtained.

d. Purple: immersing in a warm bath —60° C. (140° F.)—of 60 g. nickel ammonium sulphate, 60 g. ammonium

chloride, 1 l. water.

e. Steel-blue: a bath of 60 g. cobalt ammonium sulphate, 60 g. ammonium chloride, 1 l. water.

3. For Tin

Tin, before coloring, is either copperor brass-plated and then treated as given for these metals.

4. For Aluminum

Aluminum can generally be colored black only, either by burning in a layer of carbon produced by linseed oil or albumen, or by immersing in a 5% platinum chloride solution in water or 1% platinum chloride solution in alcohol, and left to dry in 150° C. (302° F.). The methods used for black-coloring of copper can also be applied.

5. For Iron

Black can be obtained by burning in linseed oil, tallow or wax at 400° C. (752° F.) in rotating drums, or in aqueous solution of 2% copper chloride, 2% bismuth chloride, 4% mercury chloride, 12% hydrochloric acid and 10% alcohol; the object is boiled in this solution. Iron can be burnished at 100° C. (212° F.) in a solution of 1% ferrous chloride, or 7% ferrous chloride and 0-2% mercury chloride with addition of a few drops of hydrochloric acid. A reddish-brown is obtained by applying a solution of 15 g, ferric chloride in 1 l. water and leaving it in for a few hours.

For Silver

Black is obtained by either a 1% aqueous solution of ammonium sulphide or a 5% solution of ferric chloride and rinsing in 2% caustic soda.

7. For Gold

A red-gold tint is produced by a warm solution of 115 parts salt, 230 parts saltpeter, 170 parts hydrochloric acid and 150 parts water; or of 3 parts hydrochloric acid, 1 part nitric acid, 2 parts salt in 40 parts water.

8. For Nickel

Treating with platinum chloride or sal ammoniae containing ammonium sulphide gives black and gray tints.

Black Finishing Chromium Plate U. S. Patent 1,937,629

Immerse articles for 20-30 minutes in:
Sodium Cyanide 45 lb.
Soda Ash 35 lb.
Salt 20 lb.

at temperature of 700-900° C.

Coloring Copper a Green-Blue

A malachite coating is formed on a copper anode in an aqueous solution of an alkali carbonate (8% sodium bicarbonate), using a c.d. of 1-20 amp./sq. dm. The coating may be applied to copper roofs, etc., by means of a cloth-covered roller soaked in the electrolyte. The coating is green and adherent, and changes to brochantite within a year without flaking.

Coloring Brass Cheap Rose Gold Finish

The work which must be brass is placed in the following dip until a smut is produced:

Copper Sulphate 16 oz. Muriatic Acid ½ gal. Water 1 gal.

Dissolve the copper sulphate in the water and then add the acid. The work should have a deep red smut which should be lightened somewhat by placing in a saturated salt solution for a few seconds. Plate in the regular fine gold solution, then relieve the high lights with bicarbonate of soda, replate in gold solution for a few seconds, dry and lacquer.

Blue Black Color

Copper Carbonate	1 lb.
Ammonium Hydroxide	1 qt.
Water	3 qt.

Add the water after the copper carbonate and the ammonia have been thoroughly mixed. Use at a temperature of 175° F. and immerse the work until the color is obtained (usually from ½ to 1 minute). There must be excess copper carbonate.

Verde Finishes Formula No. 1

White Arsenic	8	oz.	
Muriatic Acid	1	qt.	
Copper Acetate	2	lb.	
Copper Carbonate	1/2	lb.	
Ammonium Chloride	2	lb.	
Water	2	gal.	

Dissolve the arsenic in the muriatic acid with the aid of heat and then add the copper carbonate. Dissolve the copper acetate and the ammonium chloride in the water and mix the two solutions thoroughly. This is used with a brush. If desired as an immersion, reduce to twice the volume with water.

No. 2

Copper Acetate	4	oz.
Copper Nitrate	4	oz.
Ammonium Chloride	4	oz.
Water	1	gal.
No. 3		
Copper Nitrate	8	oz.
Ammonium Chloride	4	OZ.
Acetic Acid	4	oz.

1 oz.

1 gal.

Chromic Acid

Water

Apply lightly with brush and let dry. If finish is not even, brush again with the verde solution and let dry.

Verde Color (Tiffany Green)

Copper Sulphate	8 oz.
Ammonium Chloride	4 oz.
Sodium Chloride	4 oz.
Zinc Chloride	1 oz.
Acetic Acid	2 oz.
Water	1 gal.

The addition of 1 oz. of glycerin will prevent the green from drying too fast and produce a more even color. This solution is used for immersion and if the color is not uniform, repeat immersion as many times as desired, allowing the work to dry thoroughly between immersions.

Electrolytic Verde Finish

Potassium Bichromate	8	OZ.
Copper Sulphate	12	OZ.
Water	1	gal.

Use solution at a temperature of 80° F.; lead anodes and 8 to 10 volts. Then set color in an alkaline solution.

Brown on Brass Formula No. 1

- 01111010 - 101 II	
Golden Sulphuret of	
Antimony	4 oz.
Caustic Soda	8 oz.
Water	1 gal.

Use as near the boiling point as pos-

Scratch brush dry. If the color is not dark enough, pass through a dip composed of 2 oz. sulphuric acid, water 1 gal.

	INO.	3		
"Liquid" Water	Sulphur		-	oz. gal.

The work is immersed in this solution for a minute or so and then without rinsing immersed into a solution made of sulphuric acid 1 oz., nitric acid 1 oz., water 1 gal. If color is not dark enough, repeat both dipping operations and scratch brush dry.

Blue Color on Brass

Hyposulphite of Soda	8	oz.
Lead Acetate	4	oz.
Water	1	gal.

Use at boiling temperature and immerse just long enough to produce blue color.

Green Color on Brass

Nitrate of Iron	2 oz.
Hyposulphite of Soda	8 oz.
Water	1 gal.
TT 1 174	•

Use boiling temperature.

Verde Color on Brass

Copper Nitrate	16	oz.
Ammonium Chloride	. 4	oz.
Acetic Acid	1	qt.
Water	3	qt.

Immerse the work and let dry. If color is not uniform use a painter's sash brush which is moistened with the solution and stipple lightly.

Old English Finish on Brass

Two solutions are necessary to produce this finish, one a sulphur solution the other an acid solution.

Formula No. 1

	Sulphur		1/2 OZ.
Water		1	gal.
	No. 2		4.3

	110. 3	
Copper : Water	Sulphate	2 oz. 1 gal.

The work is thoroughly cleaned in an alkaline cleaning solution, then dipped in No. 1 solution, and without rinsing dipped in No. 2 solution. These dips are only momentary. Rinse in clean cold water and repeat dipping operations until a light color is produced.

For an even finish, scratch brush, dry and repeat dipping operations in solutions No. 1 and No. 2; finally scratch brush dry and lacquer.

Coloring Brass or Copper (Use Brush or Immersion)

Black Potassium Sulphide 2 oz. Ammonium Chloride 2 lb. Water 1 gal. Brown Ammonium Sulphide 2 oz. 1 gal. Blue Green (180° F.) Sodium Thiosulphate 1 oz. Iron Pernitrate 8 oz. Water 1 gal. Rust Brown Barium Sulphide 2 oz. Water 1 gal. Red (120° F.) Copper Sulphate 4 oz.

2 lb.

1 gal.

Salt

Water

Verde Green (75° F.)

	,
Copper Nitrate	5 oz.
Ammonium Chloride	5 oz.
Chloride of Lime	5 oz.
Water	1 gal.

Coloring Bronze

Formula No. 1

Use a boiling or near-boiling solution containing 50 to 60 g. copper sulphate per liter of water. Additions of alum (potassium aluminum sulphate) give colors tending toward the violet-red. About 20 g./l. are recommended.

Additions of verdigris give olive-green colors. About 30 g./l. are recommended, with further additions of 5 to 10 g./l. if desired.

A very pretty red may be obtained from the following:

Copper Sulphate	62.5 g.
Verdigris	10 g.
Alum	25 g.
Water	1 Ĭ.
Acetic Acid	few drops

Exact reproduction of this color is sometimes difficult.

No. 2

Bronze may be colored in the following:

Sodium	Chlorate	50	g.
Copper	Sulphate	125	
Water	-	1	ĭ.

If copper nitrate is used instead of copper sulphate, less sludge is obtained.

148 g. of copper nitrate should be used.

The following colors are obtained:

Solution near boiling—greenish gold-brown obtained in 5 minutes.

Solution near boiling—gold brown obtained in 10 minutes.

Solution cold—yellow brown obtained overnight.

The effects of additions are as follows:

Addition of ferrous sulphate—slight change toward olive green.

Addition of ferric ammonium sulphate—similar to above but lighter in color.

Addition of ferric sulphate—similar to above but with strong etching.

Addition of nickel sulphate—increase in yellow brown.

Addition of ammonium sulphate—lighter color and more yellowish brown, partly toward greenish.

No. 3

Antique Green-Oxidized Effect

After cleaning, dip and/or brush with stippling effect, using the following solution:

Water	1 gal
Iron Chloride	3 oz.
Sal Ammoniac	16 oz,
Verdigris Powder	8 oz.
Common Salt	10 oz.
Cream of Tartar	4 oz.

No. 4

If bronze is being exposed to the atmosphere, rub it with cotton waste soaked in boiled linseed oil to obtain, on aging, a dark brown adherent color.

No. 5

For brown, reddish bronze, or blue-black tones use:

Water	1 gal.
Liver of Sulphur	2 oz.
Caustic Soda	3 oz.

Use a temperature of 160° to 180° F. The time of exposure to the solution determines the color.

Coloring of Copper

The pieces to be colored are first cleaned of all oil and grease with gasoline and then lightly etched in the following solution:

Water			90	oz.
Concentrated	Sulphuric	Acid	10	oz.

They are then thoroughly washed in water before immersion in one of the following coloring solutions.

Brown to Steel Blue Color

Liver	\mathbf{of}	Sulphur	2	g.
Salt				g.
Water			100	

This bath works better when kept warm. The pieces are left in the bath until the desired color has been obtained.

Grav-Brown Color

Iron Chloride	3	g.
Water	100	
The pieces are heated and	dipped.	

Brown Color

Powdered Copper Sulphate	100 g.
Zinc Chloride	100 g.
Water	200 g.

This forms a paste which is smeared over the surfaces to be colored and allowed to dry.

Other Brown Coloring	Solutions
Liver of Sulphur	5 g.
Carbonate of Ammonia	10 g.
Water	250 g.
Copper Acetate	10 g.
Ammonium Chloride	5 g.
Ammonia (10%)	25 g.
Vinegar	160 g.
This is brushed on.	

Old copper effects are obtained by brushing sulphuric acid in the depressions and thoroughly washing off after the desired amount of green oxide has been formed.

After the colored pieces have been thoroughly washed and dried they should be polished and given a preservative coat of a suitable lacquer or the following mixture:

Carnauba Wax	100	g.
Japan Wax	100	g.
French Turpentine	1000	ğ.

Coloring Copper Formula No. 1

Potassium Chlorate 1 oz. Copper Sulphate 4 oz. Water 1 gal.

Use hot, scratch brush wet. If color is uneven, repeat coloring operation and scratch brush dry.

No. 2

A darker or more red color is produced in this solution.

Copper Sulphate	4 oz.
Nickel Sulphate	2 oz.
Potassium Chlorate	1 oz.
Water	1 gal.

Finishing operations are the same as above.

No. 3

Various shades of bronze from a chocolate color to a black can be produced in this solution.

Potassium Sulphide ½ to 1 oz. Water 1 gal.

For the light shades use cold and a short time of immersion. For darker, use hot, with longer immersion.

No. 4

Various colors are produced in any of the following solutions used either hot or cold.

Yellow Barium Sulphide	1 oz.
Water	1 gal.
No. 5	
Yellow Barium Sulphide	1 oz.
Calcium Sulphide	½ fl. oz.
Water	1 gal.

No.	(
	_

Golden Sulphuret of	4. 7	
Antimony ½	to T	oz.
Caustic Soda 1	to 1 to 2	oz.
Water		gal.
No. 7		_
Copper Sulphate	12	oz.
Acetic Acid		OZ.
Caustic Soda		OZ.
Water	1	gal.
No. 8		
Copper Sulphate	4	oz.
Copper Acetate	2	oz.
Potassium Chloride		OZ.
Water	1	gal.
No. 9		_
Copper Sulphate	8	oz.
Potassium Permanganate	1	oz.
Water	1	gal.
		-

Coloring Silver

Formula No. 1 Sulphide Coloring

Dip in solutions of sodium or potassium sulphide.

No. 2

Tellurium Black

Dissolve 1 oz. of pure tellurium dioxide in 16 oz. concentrated hydrochloric acid to which have been added 8 oz. water. Boiling the solution will probably be necessary.

The solution so obtained should be diluted with water, the amount depending on the anticipated use. For brushing, use about 1 part of the above with 2 parts water. For dipping, a much weaker solution is advisable.

Better results are obtained from a hot than from a cold solution.

No. 3

Platinum Black

Silver placed in hot 5% platinic chloride solution rapidly turns jet black.

No. 4

Iron Oxide Finish on Silver

Immerse the silver for about 5 seconds in a solution containing 1200 g. ferric chloride per l. water.

Rinse the article and immerse for 15 seconds in a solution containing 20 g. caustic soda per l. water.

Better results are obtained if the article is made the cathode in the latter solution.

No. 5 Black Nickel

For relief designs on silver, black nickel is often used. The presence of

zinc or copper in a nickel plating solution will cause distinct darkening of the aickel deposit. A simple formula is:

Water 1 l.
Nickel Ammonium Sulphate 50 g.
Ammonium Thiocyanate 10 g.
Zinc Sulphate 6 g.

Carbon anodes are used, and the silver article is made the cathode at about 3 amperes per sq. ft. Excess black nickel is removed with a tampico wheel and pumice.

No. 6

Pink Color on Silver

A pink color may be given silver by immersing it in a hot solution of copper chloride.

Antique Silver Finish Formula No. 1

Roughen surface (as by acid dipping) and then dip into the following solution:

Lead Acetate 3 g. Sodium Thiosulphate 140 g. Water 1 l. Temperature 140° F.

No. 2

Dip article into following solution:

Ortho Arsenic Acid 50 g.
Sodium Carbonate 20 g.
Potassium Cyanide 25 g.
Water 1 l.

Add the chemicals to the water in the above order, with thorough mixing of each.

No. 3

Dip article into solution containing 15 g. potassium sulphide per l. of water. Rinse in water and dip into following:

Copper Sulphate 9 g. Sulphuric Acid (Conc.) 3 g. Water 1 l.

Polish article with fine pumice and dip into weak solution of potassium cyanide containing sodium hydroxide.

Imitation Antique Silver Finish

An imitation antique silver appearance may be given iron, for example, by first cadmium plating it, and then dipping it in the following:

Potassium Chlorate	60 g.
Cupric Nitrate	40 g. 1 l.
Water	1 I.

Preventing Flaking in Steel

Flakes, especially in steels of the S.A.E. 3312 type, can be avoided by

thoroughly deoxidizing before adding the iron alloys, by mixing the bath well, by pouring at 1420-50°, by slow cooling and heating in the range 300-700°, and by forging at high temperature.

Coating Iron with Aluminum British Patent 432,212

Iron wire is exposed to ammonium chloride vapors at 500-700° C. and passed directly into a bath of molten aluminum.

Phosphate Coating for Steel Canadian Patent 351,060

Sodium Nitrate	100	lb.
Manganese Acid Phosphate	115	lb.
Copper Carbonate	19	
Water	400	gal

Coating Steel with Zinc Phosphate U. S. Patent 1,926,265

Dip steel in:

Zinc Cyanide 3 lb.

Zinc Acid Phosphate 15 lb.

Water 100 lb.

while heated at 75° C.

Foundry Parting Powder British Patent 412,931

Kieselguhr 92-97.5 lb., wax 6-2 lb. and resin 2-0.5 lb, the kieselguhr being thoroughly mixed with the molten wax and, after cooling, the mixture being ground with the powdered resin.

Improving Malleable Iron Castings U. S. Patent 2,024,014

The process for the heat treatment of malleable iron castings containing 0.6 to 5% copper comprises heating the malleabilized castings to a temperature in the range of approximately 700 to 850° C.; cooling at a rate greater than approximately 25° C. per hour to a temperature in the range of approximately 400 to 600° C.; and without further cooling maintaining in that temperature range for sufficient time to produce a substantial increase in hardness.

Increasing Carbon Content of Iron U. S. Patent 2,021,159

Add to molten metal after leaving cupola a mixture of:

Sodium Nitrate 20 lb. Carbonaceous Material 80 lb.

Case Hardening Composition

Formula No. 1 U. S. Patent 2,002,180

0. 2. 1 40010 2,002,100	
Sodium Cyanide	9 lb.
Barium Chloride	6 lb.
Barium Carbonate	8 lb.
Calcium Fluoride	2 lb.

No. 2

U. S. Patent 1,952,090

Calcium Chloride	20	lb.
Salt	10	lb.
Sodium Cyanide	0.15 - 0.3	lb.

No. 3

U. S. Patent 1,942,937

Heat metal at 1010-1065° C. in a mixture of:

Charcoal Powder	40 1	b.
Hardwood Sawdust	24 l	b.
Manganese	20 1	b.
Chromium	5 I	b.
Borax	8 1	b.
Chopped Pea Plants	3 1	b.
allowing free access of air.		

No. 4

British Patent 412,173

Metal is dipped in following:

T.T.		
Ground Rice	31 lb.	
Barium Carbonate	21 lb.	
Caustic Soda	1 lb.	
Glucose	5 lb.	
Silica	3 lb.	
Water	39 lb.	

After drying the coated metal, heat to $900-950^{\circ}$ C. in a non-oxidizing atmosphere.

Hardening Steel Formula No. 1

Austrian Patent 142,401

Potassium Ferrocyanide	70-80	kg.
Soda Ash	2-5	kg.
Salt	6-12	kg.
Acetylene Carbon	3-8	kg.
Potassium Carbonate	2-3	kg.
Ammonium Chloride	2-3	kg.
Gum Arabic	2-3	kg.

The above mixture is strewn over the steel which is then heated.

No. 2

U. S. Patent 2,016,477

Soybean Powder	00	lb.
Boybean Fowder	90	TD.
Sodium Cyanide	3	lb.

Salt Soda Ash Ammonium Chloride Barium Carbonate Potassium Dichromate	0.8 0.2 4 1	
Potassium Dichromate	1	lb.

No. 3

British Patent 416,179

Coat with following and heat to carburizing temperature:

urizing temperature:	
Carbon Powder	40 lb.
Barium Carbonate	20 lb.
Nickel Steel (20%)	
Turnings	15 lb.
Asbestos Fiber	12 lb.
Sodium Silicate (d. 1.33)	13 lb.

No. 4 Patented

Immerse in a fused salt bath of:

Calcium Cyanide	15-40 lb.
Sodium Nitrate	20-40 lb.
Barium Carbonate	10-15 lb.
Salt	5-10 lb.

Temperature is maintained at 760-960° C. and a current of ammonia gas is passed through the bath to produce a nitride case.

Air Hardening Steel U. S. Patent 1,976,341

An air quenched article of alloy steel is composed of about 3 to 4% copper, about 0.1 to 0.25% carbon, and about 1.5 to 2% manganese, the balance being substantially all iron.

Hydrogen Chloride Resistant Steel German Patent 596.023

Copper	43-74	kg.
Nickel	10-25	kg.
Zinc	3.5 - 14.5	kg.
Tantalum	0.5 - 7	kg.
Manganese	0.3 - 1.5	kg.
Bismuth	0.2 - 8.5	kg.
Molybdenum	0.4-7	kg.
Silver	0.1 - 4.5	kø.

Surface Carbonization of Steel U. S. Patent 1,950,116

Etch surface in 15% nitric acid; wash; dry; heat at 900° C. in a hydrocarbon vapor.

Anti-Carburizing Composition U. S. Patent 1.982.718

		 -	,			
Copper Chloride	Э			2	lb.	
Oxalic Acid				3	lb.	

Lead Oxide	1	lb.
Copper Sulphate	5½	lb.
Water	5	lb.
water	อ	ın.

Metallographic Etching Agent
Copper Ammonium Chloride 3 g.
Hydrochloric Acid 50 cc.
Ferric Chloride 15 g.
Water 25 cc.

Etching Hardened Steel

Mercuric Nitrate	5	oz.
Nitric Acid	38.5	oz.
Water	89.5	oz.

Etching Stainless Steel Formula No. 1

Nitrie Acid	32	oz.
Hydrochloric Acid	3	oz.
Denatured Alcohol		oz.
Water	96	oz.
Solution used cold.		

No 2

X101 M	
Ferric Chloride	20 g.
Hydrochloric Acid	20 g.
Water	60 cc.

This solution may be used warm at 120° F. or electrolytically.

Steel Pickling Inhibitor U. S. Patent 1,932,015

Di-o-tolylthiourea	4	lb.
Evaporated Waste Sulphite		
Liquor	6	lb.
Salt	10	lb.
Soda Ash	. 1	lb.
The above is formed into ble	ocks.	

Metal Pickling Inhibitor Canadian Patent 353,320

Pyridine	80	g.
Benzyl Chloride	140	

Heat to $160-170^{\circ}$ C. and cool to $75-100^{\circ}$ C. and then dilute with any solvent.

Ore Briquettes for Open Hearth Furnaces

Ore	100 lb.
Cast Iron Shavings	10 lb.
Salt	1 lb.

More satisfactory results are gotten by using above briquettes than when using dust ore.

Age Hardening Silver U. S. Patent 1,984,225

Sterling silver capable of age hardening to a hardness of from 84 Rockwell B to 94 Rockwell B consists of pure silver at least 92.5%, copper 2.5 to 7.4% and aluminum 0.1 to 5%.

A process of making sterling silver articles of a hardness of from 80 Rockwell B to 94 Rockwell B consists in first alloying at least 92.5% silver, from 7.4 to 2.5% copper and from 0.1 to 5% of a metal selected from the group consisting of aluminum, magnesium, lead, antimony, and beryllium, then fabricating the article to form by known cold working operations, then subjecting the article to a preliminary anneal and quench from about 1150° F. to 1400° F. and finally subjecting the article to an age hardening heat of about 570° F. for about one hour.

PHYSICAL PROPERTIES OF METALS

			Melting	Point	Weight
	Specific	Specific	Deg. Cen-	Deg. Fah-	
Metal	Gravity	$\mathbf{H}\mathbf{eat}$	tigrade	renheit	Cubic Inch
Aluminum:					
(Cast)	2.56	.2185	658	1217	.0924
(Rolled)	2.71				.0978
No. 38 Alloy (Rolled)	2.74				.0989
No. 12 Alloy (Rolled)	2.82	• • • •	624	1156	.1018
Antimony	6.71	.051	630	1166	.2424
Bismuth	9.80	.031	271	520	.3540
Brass	8.51	.094			.3075
Cadmium	8.60	.057	321	610	.3107
Calcium	1.57	1.70	810	1490	.0567
Chromium	6.80	.120	1510	2750	.2457

PHYSICAL PROPERTIES OF METALS—Continued

Metal	Specific Gravity	Specific Heat	Melting Deg. Cen- tigrade		Weight in Lbs. per Cubic Inch
Cobalt	8.50	.110	1490	2714	.3071
Copper	8.89	.094	1083	1982	.3212
Gold	19.32	.032	1063	1945	.6979
Iridium	22.42	.033	2300	4170	.8099
Iron	7.86	.110	1520	2768	.2634
Iron (Cast)	7.218	.1298	1375	2507	.2605
Iron (Wrought)	7.70	.1138	1500-1600	2732 - 2912	.2779
Lead	11.37	.031	327	621	.4108
Lithium	0.57	.941	186	367	.0213
Magnesium	1.74	.250	651	1204	.0629
Manganese	8.00	.120	1225	2237	.2890
Mercury	13.59	.032	-38.7	-37.7	.4909
Monel Metal	8.87	.127	1360	2480	.320
Nickel	8.80	.130	1452	2646	.319
Platinum	21.50	.033	1755	3191	.7767
Potassium	0.87	1.70	62	144	.0314
Silver	10.53	.056	961	1761	.3805
Sodium	0.97	.290	97	207	.0350
Steel	7.858	.1175	1330-1378	2372-2532	.2839
Strontium	2.54	.074	• • • •	• • • •	.0918
Tantalum	10.80	• • • •	2850	5160	.3902
Tin	7.29	.056	232	450	.2634
Titanium	5.3	.130	1900	3450	.1915
Tungsten	19.10	.033	3000	5432	.6900
Uranium	18.70	• • • •	••••	• • • •	.6755
Vanadium	5.50	• • • •	1730	3146	.1987
Zinc	7.19	.094	419	786	.2598

Protecting Aluminum from Corrosion Immerse for 10 minutes in bath of following at 50-60° C.

Formula No. 1

Sal Soda	125	g.
Sodium Chromate	8	g.
Ammonia	25	cc.
Water	1	1.

No. 2

Anodic treatment at 12 volts for 5 minutes and 15 volts for 5 minutes in following bath:

Oxalic Acid	25	g.
Sodium Chromate	17	g.
Sodium Dihydrogen Sulphate	3	ğ.

Hardening Aluminum U. S. Patent 1,930,463

Pack in a mixture of:	
Magnesium	95 lb.
Magnesium Oxide	5 lb.

and heat at 420° C. in an atmosphere of carbon dioxide until the magnesium diffuses into the surface of the aluminum.

Non-Seizing Aluminum U. S. Patent 1,978,112

Dip the aluminum in a bath of molten aluminum stearate.

Rustproofing Iron U. S. Patent 1,949,921

Phosphoric Acid	(85%)	20	fl.	oz.
Ethyl Alcohol		20	fl.	oz.
Water		30	fl.	oz.
Isopropyl Ether	0.	7 - 3.5	ff .	OZ

Radiator "Rust" Preventative U. S. Patent 1,940,041

Borax	36	lb.
Sodium Salicylate	30	
Sodium Nitrate	7	lb.
Use 73 grains per	quart of water	

Corrosion Inhibitor

20	lb.
15	lb.
50	lb.
2	lb.
อ	lb.
100	lb.
	15 50 2 5

Non-Corrosive (Ethyl) Alcohol U. S. Patent 1,927,842

About 0.03 per cent of sodium carbonate or the equivalent of sodium acetate, borax, sodium lactate, or the corresponding potassium salts, is added to commercial alcohol to give pH 7, thereby preventing corrosion of the metal containers.

Non-Corrosive Zinc Conduit Alloy German Patent 614,996

Zinc	83-95	kg.
Aluminum	13 - 3	kg.
Manganese	1-2	kg.
Cadmium or Silicon	3-0	kg.

Silver Tarnish Prevention British Patent 430,795

A jar containing the following is placed in display cases containing silver:

Calcium Chloride,		
Granular	88-94.9	g.
Copper Sulphate,		_
Anhydrous	5-10	g.
Talc	0.1 - 2	g.

Removing Rust from Iron Formula No. 1

Soaking 12 hours in Petroleum

Doubling 12 Hours in 1 Co.	orcuii.	L
No. 2		
Make up:		
Spindle Oil	65	g.
Paraffin Scales or Ceresin,		
Yellow Pumice Powder	$\frac{15}{20}$	g.
	20	g.
No. 3		
Dissolve:		
Water Stannous Chloride	1000	g.
Mercuric Chloride	10	g. g.
		8.

Caustic Soda 10 g. Zinc Powder 10 g.

Use:

No. 4

Removing Rust From Tools

By using a solution of ammonium citrate, rust may be completely removed

from tools. If the solution is used warm, then one or two hours will suffice, but if used cold, it is best to allow the tools to remain in the liquid overnight. A tablespoonful of the ammonium citrate crystals may be used to a pint of water, although the proportions are not important. The solution will serve repeatedly until depleted.

For tools of awkward shape such as try-squares and large steel squares, a cardboard mailing container may be used in place of a vat, crock, or other container, if it is first impregnated with hot

paraffin wax.

Rust and Oil Remover U. S. Patent 1,935,911

Brush with:		
Phosphoric Acid (75%)	69.5	lb.
Butyl "Cellosolve"	17	lb.
Oleic Acid	0.5	lb.
Saponin	1	lb.
Water	12	lb.

Cleaning Motor Nameplates

Cleaning tarnish, grease and dirt off the nameplate of motors and generators in order to read the figures and other data is facilitated by the use of a wad of crinkled tin foil. The nameplate is not scratched or marred by this material as is the case when an abrasive is used for removing the accumulated dirt.

Decarbonizing Lining for Cast Iron Molds

Russian Patent 35,331

Brown Iron Ore	68	lh
Refractory Clay	30	
Potassium Permanganate	2	lb.

Soldering Fluxes for Iron and Non-Ferrous Metals

Stainless Steel

Water

Borax	75-25	
Boric Acid	25-75	oz.

Make into paste with alcohol. Galvanized Iron

Hydrochloric Acid	750	cc.
Water	250	cc.
Zinc add until no more will	disso	lve

men add a	solution of	
Ammonium Water	Chloride	50 g.
water		170 ec.

170 cc.

then add	following	solution:	
Stannous			30 g.

To form a paste solder of this type work in potato starch to desired consistency.

Aluminum Sheets

Formula No. 1		
Rosin	2	lb.
Tallow, Ox	2	lb.
Zinc Chloride	1	lb.
No. 2		
Olive Oil	50	lb.
Tallow	40	lb.
Rosin, Powdered	25	lb.
Saturated Ammonium Chlo-		
ride Solution	$12\frac{1}{2}$	lb.
l'in		
Rosin, Powdered	1	lb.
Tallow	2	lb.
Olive Oil	2	lb.
Saturated Ammonium Chlo-		
ride Solution	2	lb.

Aluminum Solder Formula No. 1

Tin	76 oz.
Zine	20 oz.
Aluminum	3 oz.
Antimony	0.6 oz.
Lead	0.2 oz.
Copper	0.2 oz.

No. 2

French Patent 775,492

The solder contains cadmium, lead and zinc in the proportions of 4, 4, 3 and 2-10% of zinc chloride.

No. 3

French Patent 776,958	Frenc	h Paten	t 770	3,958
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Zine	89-95		lb.
Aluminum	10-4		lb.
Silicon	0	.4	lb.
Iron	0	.4	lb.
Zirconium	0	.2	lb.
N	o. 4		

British Patent 426.526

Zinc	22	lb.
Tin	14	lb.
Mercury	3	lb.
Aluminum	11/2-1	lb.
Lead	1/2-1	lb.

Aluminum Soldering Fluxes British Patent 413,141

Claim is made for fluxes containing cadmium chloride and stannous bromide, preferably in admixture with more than one of the following: cadmium iodide, ammonium chloride, zinc chloride, zinc bromide, fluorides, chlorodiphenyl, p-dichlorbenzene. A preferred composition

is stannous bromide 28, cadmium chloride 20, cadmium iodide 10, ammonium chloride 25, ammonium fluoride 2, zinc chloride or zinc bromide 5%; 4 parts of this mixture are made into a paste with 6 parts of chlorodiphenyl and/or p-dichlorbenzene.

Soldering Iron Tip Alloy British Patent 431,637

Copper	97	lb.
Cobalt	2.6	lb.
Bervllium	0.4	lb.

Heat this for one hour at 900° C. Quench in water, reheat to 500° C. for one to two hours and allow to cool.

Cast Iron Soldering

Add to muriatic acid, zinc sufficient to "kill" it, and drop in several small pieces copper before action ceases. Use this solution on cast iron that has been filed bright, and solder in the usual way.

Hard Solder for Cast Iron Copper 60 lb. Zinc 40 lb. Tin 1 lb. Iron 1 lb.

Chain Link Solder

U. S. Patent 2,003,865

Tin	1 1	b.
Copper	2 1	b.
Borax	3 1	b.

Hard Solders

	marcor Ct	Coldela			
German Silv	er and	Nickel			
Silver					lb.
Copper Tin					lb.
Thin Copper				Ü	10.
Silver				65	Ib.
Copper				24	lb.
Zinc				11	lb.
Heavy Brass					
Silver				30	lb.
(Camman			40	-0	77

Copper 40-50 lb. Zinc 20-30 lb.

Austentic Stainless Steels

Silver				10	lb.
Copper				50-60	lb.
Nickel	1.2			3	lb.
Zinc				37-27	lb.

Soft Soldering Monel and Nickel

Monel metal and nickel are soft soldered readily. Many of the soft solders regularly used in the copper shop will make suitable joints in both of these metals. In making a lock seamed joint, for example, it is definitely recommended that the edges of the sheet be tinned, that is, coated with a thin film of soft solder before forming the sheet and before lock seaming.

Once the sheet has been properly tinned, it is then very easy to flow in the soft solder and make a tight joint which

is reasonably strong.

Similarly, in sweating a tube into a header, if both the header and end of tube are tinned first, then assembled, heated, and solder flowed in, a sound joint will be obtained. It is necessary in all soldering work to have surfaces clean and bright if joints are to hold.

It must be remembered that the strength and ductility of soft solder is not of a very high order and for that reason soft solder is not recommended where considerable vibration is apt to be

involved.

Soldering Flux for Stainless Steel U. S. Patent 1,968,841

Boric acid, three parts; borax, two to three parts; and ammonium chloride, one and one-half to three parts, together with a liquid from the group consisting of water and hydrogen peroxide, in quantity to make a thick paste.

Soldering Flux for Stainles	s Steel
Zinc Chloride	37 oz.
Acetic Acid	23 oz.
Hydrochloric Acid	40 oz.

Tin Plate Solder

Ammonium Chloride	4 oz.
Zinc Chloride	48 oz.
Hydrochloric Acid	1 oz.
Water	47 oz.
Dilute to required strength	with water

Crankshaft Heat Treatment
Shafts are heat treated in gas fired
furnaces as follows:

Heat to 1650° F., hold for 20 minutes. Air quench to a minimum of 1200° F. Reheat to 1480° F., and hold 1 hour. Cool in the furnace to 1000° F. in another hour.

The alloy for casting is melted in four 15-ton electric furnaces according to the latest approved practice. The charge is

made up of approximately 50 per cent return shop scrap (gates and risers) and 50 per cent steel scrap.

Drawhead Casting Heat Treatment

To obtain the best combination of mechanical properties, the castings are given a simple heat treatment, as follows:

Heat to about 1650° F., hold at heat 1 to 1½ hours per inch thickness of heaviest section, and cool in still air to a black heat. Reheat to 1200-1250° F., hold at least 1 hour per inch and cool in air or furnace. This treatment is not difficult and can be performed very readily with ordinary equipment. Usually the cost of such a treatment is no greater than that for simple annealing.

Oil Well Tool Heat Treatment

The heat treatment of these steels (used either for slip socket or tool joint) is quite similar. After forging it is advisable to anneal the steel to relieve any forging strains and at the same time put it in a readily machinable condition. One of the simplest treatments for doing this is to heat the steel to above 1600/1650° F. and cool in the furnace until black or, if removed from the furnace, pack in lime or ashes so that it cools slowly. The forging should then be machined and the final heat treatment performed as follows:

Heat to about 1550° F.; hold at this temperature until heated through thoroughly; quench in oil. The tempering operation will depend upon the hardness specifications. This steel is quite tough in the hardness range of 280/320 Brinell which could be secured by using a drawing temperature around 900° F., holding at this temperature until heated through thoroughly in the heavy sections. While final machining can be performed in this hardness range, it must be done very slowly, and it is desirable to use a lower hardness range, such as about 240/280 Brinell which is obtained with about 950° F. draw. The physical properties secured at these hardnesses is about as follows:

Tensile Strength Yield Point	145,000 120,000	
Elongation in 2" Reduction of Area	18% 57%	Picin

For the slips a case hardened steel such as S.A.E. 2315 is used, arrangements to be made so that only the teeth are case hardened. This can be accomplished

by copper plating the piece before cutting the teeth so that the copper remains on all the parts except the teeth. Use a case hardening temperature of 1650/1700° F., cooling in the box and reheating the parts to a temperature of 1475° F., quenching in oil, and tempering by heating to 275-300° F. This treatment will toughen the core of the part so that it will be sufficiently hard not to stick or gall against the socket.

Heat Treatment of High Strength Shafting

Heat treatment for S.A.E. 3340: Oil quench from 1500° F. and temper at 800° to 900° F.

Heat treatment for Ni-Cr-Mo: Oil quench from 1575° F., temper at 900° to 1000° F.

Brake Drum Heat Treatment

The heat treatment given brake drums is heating to 1600° F., holding there for 30 min., then cooling rapidly in the furnace to 1450° F., followed by cooling in 2 hours to 1350° F. and then in 1 hour to 1000° F.

Valve Gear Metal Heat Treatment

A nickel-molybdenum case-hardening steel corresponding to S.A.E. composition 4615 is used. This material can be machined to the finished size in a soft state and then should be carburized by the pack method, at a temperature between 1650 and 1700° F., until a case about 1/30 in. in depth is secured. For the best results we would recommend quenching from the carburizing box into oil. This should be followed by a reheating to a temperature of 1375 to 1400° F. and quenching in oil, then temper at about 275° F. This treatment will result in a very hard case which should show excellent wearing properties.

Carburizing Nickel Steel

(1) A simple and economical treatment where refinement of the case is not important, is to carburize at 1600° F. and quench in oil directly from the box, followed by tempering at 250 to 350° F. (2) Or, if cooled in the box after carburizing, then heat to 1475–1500° F. and oil quench, then temper as above, to get a refined and tough core which will back up the hardness of the case. (This is not recommended if the carbon content of the core is over about .18%, as brittleness may result.)

(3) Cooling in the box, oil quenching

from 1325 to 1375° F. and tempering, is recommended where a hard and refined case is the main requirement. (4) If refinement of both case and core is demanded, and economy and speed is not so important, a double treatment should be given, as follows: Carburize at 1600° F., cool in box. Quench in oil from 1500–1550° F., and again from 1325–1375°. Temper at 250–350° F. as required. This will give a very hard case and a ductile core, and is much used on gears of fine pitch.

Grinding Wheels U. S. Patent 1,937,043

Carborundum 900 g., is mixed with furfuraldehyde 10 cc. till moist then with a phenolic resin 100 g., and the mixture is pressed into shape at less than 80° C. The articles are then heated at a suitable temperature until complete hardening occurs.

Aluminum Welding Flux

Potassium Chloride 79 oz.
Salt 16 oz.
Potassium Bisulphate 5 oz.
The above is best used with welding

Bronze-Welding

aluminum containing 4% silicon.

Bronze-welding, as a general term for actual bronze-welding and for bronze-surfacing, is used today for joining metals of high melting points, as cast iron, steel, nickel, copper and their alloys, by the use of a bronze bonding material. For use with the oxy-acetylene flame, a rod of 59% copper, 40% zinc and 1% tin is generally used, while recently other elements as silicon, manganese, iron have been added. Lead is objectionable as it increases porosity of the weld metal.

Welding Rods for Copper, Steel and Bronze

U. S. Patent 2,009,977

Silicon	3.5	lb.
Tin	0.5	lb.
Phosphorus	0.05	lb.
Copper	96	lb.

Welding Zinc and Zinc Alloy Castings

The welding of zinc requires some care because of its low melting point and the tenacious character of the oxide. A gas flame should be used with welding rod of the same metal and a flux of ammonium chloride and water. The welding operation always weakens the surrounding

metal and should, if possible, be followed by a cold working operation to refine the

grain.

Zinc alloy castings containing aluminum are extremely difficult to weld and the success of the operation depends largely on the technique of the welder.

Welding Electrode Coating Canadian Patent 341,572 Formula No. 1

Shredded wood 100, sodium silicate 80, calcium carbonate 5, kaolin 5, silicomanganese 5 and peanut oil 5 parts. The coating in a plastic state is applied to the core and then baked or dried.

No. 2

U. S. Patent 1,968,984

Barium	Chloride	20-50	lb.
Lithium	Fluoride	4-6	lb.

To the above add 75-45% of following mixture.

Salt		40-50	lb.
Potassium	Chloride	60-50	lb.

No. 3

U. S. Patent 2,000,861

Slip clay 40-60 parts, iron oxide 20-30 parts, calcium carbonate 20-30 parts, feldspar 15-30 parts, rutile 5-20 parts, manganese ore 5-15 parts, carbonaceous material 5-15 parts, ferromanganese 5-20 parts, ferrochrome 2-8 parts and dextrin 1-15 parts by weight.

Welding Rod for Bearing Metals U. S. Patent 1,926,412

Zinc	90 lb.
Copper	5 lb.
Antimony	5 lb.

Welding Rod Coating Formula No. 1

Canadian Patent 347,320

Calcium Carbonate 8	lb.
Barium Carbonate 9	lb.
Titanium Dioxide 22	lb.
Calcium Fluoride 11	lb.

tion of
Potassium Silicate 2 lb.
Water 1 lb.

No. 2

Suspend above in sufficient of a solu-

U. S. Patent 1,992,792

Titanium Dioxide	1 lb.
Talc	1 lb.
Feldspar	1 lb.
Sodium Silicate	3 lb.
Water	to suit

Aircraft Engine Alloys

Use case hardened 5% nickel steel (S.A.E. No. 2512) for aircraft engine gears. The crankshafts should be forged of a nickel-chromium steel such as S.A.E. 3240, or nickel-chromium-molybdenum steel of the following approximate composition:

Carbon	0.40 - 0.50	lb.
Manganese	0.45 - 0.75	lb.
Nickel	1.50 - 2.00	lb.
Chromium	0.60 - 0.90	lb.
Molybdenum	0.15 - 0.25	lb.
Iron	to make 100	lb

Heavy Duty Axle Alloy

Carbon	0.35-0.45 lb.
Nickel	1.50-\(\pi\).00 lb.
Chromium	0.60-0.80 lb.
Manganese	0.60-0.80 lb.
Molybdenum	0.30-0.40 lb.
Iron	to make 100 lb.

Nickel Steel Pin and Bearing Alloy

5% nickel steel such as S.A.E. 2512, with the carbon at the upper end of the

range, say 0.15% is used.

Carburize this steel at 1600-1650° F. The most suitable depth of case will depend upon the dimensions of the pin, and normally should not be more than 15% of its diameter. The cooling after carburizing should preferably be done in the box, but it is recommended that it be as rapid as convenient, such as allowing the box to cool in free air or possibly in an

For the hardening operation a single quench would be advisable at a temperature just high enough to refine the core. On these small pieces a temperature around 1440-1450° F. would be sufficient. The tempering operation on this steel should be at 275° F. The complete treatment will give maximum core strength, combined with very good toughness.

air blast.

Hard Tool Steel Alloys Japanese Patent 101,748

Mold following under high pressure at 1600–1800° C.

Formula No. 1

Vanadium Powder	,	lb.
Tungsten Carbide 9	5	lb.

No. 2

Titanium Powder	5	lb.
Tungsten Carbide	95	lb.

No. 3		
Vanadium	3	lb.
Titanium	2	lb.
Tungsten Carbide	95	lb.

Steering Knuckle and Spring Bolt Alloy
A case hardened steel of the following
composition is used.

Carbon	0.12 - 0.20	lb.
Manganese	0.30 - 0.60	lb.
Nickel	3.25 - 3.75	lb.
Molybdenum	0.20-0.30	
Iron	to make 100	lb.

Punch and Die Alloys

Use	steels	co	nt	ai	nin	g

Carbon	0.6 - 0.65	lb.	0.6 - 0.65	lb.
Manganese	0.3 - 0.6	lb.	0.3 - 0.6	lb.
		lb.	1.5-2	lb.
Chromium	0.9 - 1.25	lb.	0.6 - 0.8	lb.
Molybdenu	m —		0.2 - 0.4	lb.
Iron		to	make 100	lb.

It should be thoroughly annealed after forging as follows: Heat to 1550/1575° F., air-cool, reheat to 1200/1250° F., hold for 6 to 8 hours and cool very slowly. To harden, heat to 1425° F., quench in oil, and temper for 1 hour at 425/450° F.

Shovel Dipper Teeth Alloy

Carbon	0.4-0.50	lb.
Nickel	3.0 - 3.50	lb.
Chromium	1.0 - 1.25	lb.
Molybdenum	0.3-0.40	
Iron	to make 100	lb.

Heat treatment:

Heat to 1750° F., hold 1½ hours per inch thickness; air cool. Reheat to 1250° F., hold at least one-hour per inch thickness; cool in air or furnace. Some foundries furnish the teeth in this condition, while others claim better wear by giving the tips a second treatment for hardening. This is done by heating the point or tip to a distance of 2 in. or 3 in. dependent upon the size and shape of the tooth), to a red heat (1500-1800° F.) and cooling rapidly with an air blast. If it is found that the points are too brittle the whole tooth may be drawn at 700-800° F. Sometimes this tip hardening treatment is given to the castings after a plain annealing of the whole tooth, thus eliminating original air quenching and drawing treatment described at the beginning of this paragraph.

Another steel used quite successfully for shovel teeth in this service is the following:

Carbon 0.40-0.5 lb. Nickel 1.75-2.0 lb.

Chromium	0.70 - 0.9	lb.
Iron	to make 100	lb.

These castings are given either an annealing or air quenching treatment as described above for the nickel-chromium-molybdenum steel. The tips are then reheated to a red heat and quenched in oil. The whole casting is then drawn at 700-900° F., depending upon the hardness required.

A steel which is giving excellent service in castings subjected to wear, has the following composition:

Carbon	0.35 - 0.45	lb.
Manganese	1.25 - 1.50	lb.
Nickel	2.25-2.50	lb.
Iron	to make 100	lb.

Acid Resisting Alloy Patented

Molybdenum	0.5-10	lb.
Tin	4-5	lb.
Lead	95.5-85	lb.

Antifriction Alloy British Patent 413,209

Copper	67.5	lb.
Lead	25	lb.
Tin	5	lb.
Nickel	1	lb.
Antimony	0.5	lb.
Cadmium	0.5	lb.
Zinc	0.5	lb.

Hard Aluminum Alloy British Patent 406,161

Aluminum	89 -94	lb.
Magnesium	1.5	lb.
Copper	3.7- 5.5	lb.
Nickel	0.2 - 1	lb.
Silicon	0.2 - 1	lb.
Manganese	0.4 - 2	lb.

Aluminum Alloy for Chill Casting U. S. Patent 1,997,494

Aluminum	75 -95	lb.
Iron	2 -10	lb.
Antimony	0.5 - 15	lb.
Magnesium	0.2 - 0.4	lb.

Oxidizing Nickel Silver

Hydrochloric Acid	1 -	gal.
White Arsenic	10	oz.
Copper Sulphate	10	oz.
Ferric Chloride	2 -	oz.
Copper Acetate	2	oz.
Ammonium Chloride	1	OZ.
Hyposulphate of Soda	11/2	OZ.

Heat the hydrochloric acid, and when hot put in the white arsenic. When the white arsenic is completely dissolved, mix

in the balance of the formula.

It must be definitely understood that this solution can only be used while cold. The article can be placed in a plater's basket or wired and dipped possibly half a dozen times in the solution, rinsed in cold water and then dipped in a solution of sodium cyanide and then rinsed again in cold water. After this rinse, the article should again be dipped in the oxidizing solution and the process is then complete.

The result should be a jet black oxide which can be scratch brushed if a solid black is desired, and can be readily

spotted off for highlights.

Copper Alloy Resistant to Sea Water U. S. Patent 1,956,251

Silicon	1 - 3.25	lb.
Tin	0.5 - 1.5	lb.
Iron	0.75-1.27	lb.
Lead	0 - 2	lb.
Copper	97.75-93.98	lb.

Copper Alloy Spot Welding Electrode U. S. Patent 1,957,214

Electrodes	are	tipped	with	follo	wing
Cobalt				2.6	lb.
Beryllium				0.4	lb.
Copper				97	lb.

Cold Working Copper Alloy U. S. Patent 1.936.397

Silicon	0.75	lb.
Manganese	0.25	lb.

Non-Staining Copper Alloy U. S. Patent 2,007,430

Nickel			1 to	5	lb.
Cobalt		0.2	5 to	2	lb.
Silicon	Ĺ	0.2	5 to	2	lb.
Alumi	aum		1 to	5	lb.
Molyb	denum	0.2	5 to	3	lb.
Iron		0.1	0 to	1	lb.
Calciu	m -	0.0	5 to	0.5	lb.
	r of an	amount	to o	compl	.ete
100 lb. 1	nass.			_	

High Melting Copper Alloy German Patent 597,938

Beryllium	0.3-1	0 lb.
Aluminum	0.5-1	2 lb.
Copper	99.2-8	88 lb.

Low Cost Dental Alloy

85	οz
10	oz
. 5	ΟZ
	10

Cheap Dental Inlay Alloy

	19.29	lb.
	79.29	lb.
	0.71	lb.
	0.71	lb.
•		19.29 79.29 0.71 0.71

Cast Denture Alloy Canadian Patent 342,946

Chromium	17.5	lh
Cobalt	57	lb.
Tungsten	3	lb.
Nickel	21	lb.
Iron	1	lb.
Carbon	0.5	lb.

Dental Alloy

French Patent 43.121

Gold	20-15	oz,
Copper	3-12	OZ.
Silver	65-63	oz.
Zinc	7-8	oz.

Dental Filling Alloy German Patent 603,456

Bismuth	62.5 g.
Tin	37.2 g.
Gallium	1.3 g.

Dental Alloy Casting Mold British Patent 412.303

Plaster of Paris	40 lb.
Cristobalite	45 lb.
Tridymite	10 lb.
Quartz	5 lb.

Dental and Jewelry Alloy U. S. Patent 1,965,012

	 •	
Gold	5-15	oz.
Palladium	22-30	oz.
Silver	37–50	oz.
Copper	10-20	oz.
Indium	0.5 - 5	oz.

Imitation Gold Alloy French Patent 776.806

		,	
Copper		80-82	g.
Zinc		11-15	
Tin		3-5	g.
Nickel		2	g.

During fusion add the following per 100 g. of alloy.

Cream of Tartar 9 g.
Magnesium Oxide 6 g.
Ammonium Chloride 3.5 g.
Lime 1.5 g.

Lead Calcium Alloys British Patent 412,316

Lead and pea size pieces of calcium carbide are mixed at 650-700° C. in presence of fused slag consisting of salt, calcium chloride and calcium fluoride. Alloys containing 3-3.5% calcium are obtained in 8 to 10 hours.

Lead Storage Battery Alloy British Patent 411,524

T'ellurium	0.05	lb.
Antimony	6	lb.
Lead	93.95	lb.

Non-Corrosive Magnesium Alloy German Patent 613,511

Zinc	1-10	lb.
Iron	0.02 - 1	lb.
Silver	0.05 - 3	lb.
Magnesium	98.93-86	lb.

Radium Beam Therapy Alloy

	Nickel Copper					lb.
	Tungst					lb.
_		the	powdered	metals	at	1250-
1	.350° C.					

Arc-Light Reflector Alloy German Patent 615,119

Cobalt or Nickel	20-60	lb.
Tungsten or Molybdenum	15 - 50	lb.
Chromium	30-40	lb.
Carbon or Silicon	1-5	lb.

Electric Light Reflector Alloy British Patent 412,074

A	luminum	60	lb.
\mathbf{s}	ilver	25	lb.
M	Iagnesium	15	lb.

Mirrors of Silver-Copper Alloy Canadian Patent 348,131

Prepare solution No. 1 by adding to 16 oz. of silver nitrate, 11 oz. of ammonia (26°) and, after the solution is complete, 16 oz. of distilled water; cool,

filter and add to the filtered solution an additional 144 oz. of distilled water.

Prepare solution No. 2 by dissolving 1 lb. crystallin copper sulphate in 64 oz. of distilled water, filter and place in a dark bottle.

For solution No. 3, to 64 oz. of distilled water add 2 lb. of crystallin Rochelle salt, heat to boiling and add 1 oz. of silver nitrate dissolved in 4 oz. of distilled water. To this mixture at the boiling point add 4 oz. of solution No. 2 and boil for at least 10 minutes; then cool, filter and place the filtered solution in a dark bottle.

For solution No. 4, dissolve 1 lb. of powdered tartaric acid in 48 oz. of distilled water, let stand 1 week and filter.

Prepare the final solution from distilled water, 64 oz.; solution No. 1, 2 oz.; solution No. 3, 2 oz.; and solution No. 4, 3 dr. Polish and brush with water the glass that is to be coated; then apply a weak solution of tin chloride with a felt block or bristle brush, rinse with water and lightly brush. Treat the surface with the final solution, and when the first coating of silver-copper alloy is deposited brush well to obtain a clean metallic surface. A second coating of the alloy may be applied and similarly polished. Apply a coating of shellac to the dried coated surface and then cover with paint.

Galena Blue Mirror (Non-Glaring) U. S. Patent 1,988,663

Solution No. 1

Lead Nitrate	2	oz.
Distilled Water	32	oz.

No. 2

Potassium Hydroxide,		
Sodium Hydroxide or		
Other Similar Alkali Agent	4	oz.
	32	oz.

No. 3

Thiourea	(Thiocarbamide)	2	oz.
Distilled	Water		oz.

In preparing the above solutions care must be taken to insure complete dissolution of the chemicals and each solution should be shaked well before using. In order to produce a lead sulphide film or layer upon the glass or other surface to be treated either of two processes may be employed, one being designated as the "hot" process and the other as the "cold" process.

In either process, the glass or other surface to be coated is initially block

polished or hand rubbed with rouge, after which it is well brushed with water. Following this water brushing operation, a weak solution of tin chloride is applied to the surface to be treated preferably by means of a felt block or bristle brush. The surface is then rinsed well with water and lightly brushed.

The glass so treated is then placed in a horizontal plane and accurately leveled with wedges, the surface to be coated being uppermost. In the "hot" process, after the glass has been initially treated, washed and leveled as just described, the following Solution No. 4 is poured upon the surface to be coated:

No. 4

Distilled Water	4 oz.
Solution No. 1	1 oz.
Solution No. 2	1 oz.
Solution No. 3	1 oz.

Attention is here directed to the fact that in preparing Solution No. 4, the numbered solutions are added to the distilled water in the order given above and that Solution No. 3 is not added until just before the final solution is to be poured upon the glass. Following the application of the tin chloride solution the surface to be treated must be kept wet until the final solution has been applied thereto. As much of the final Solution No. 4 is poured upon the leveled surface as the latter will hold without the solution running over the edges. Heat is uniformly applied to the glass preferably by placing the glass upon a table or bed the surface of which is heated to the required temperature.

In a relatively short time (about 15 minutes) lead sulphide will have deposited out of the final solution and upon the glass. The excess solution is then removed from the glass surface, preferably with a piece of chamois, after which the deposited film is well wiped to obtain a clean metallic surface. second application of the final Solution No. 4 is then made. In about 10 minutes a second coating or film of lead sulphide will have deposited out of solution upon the first coating, the second coating being also wiped and dried with the chamois. When deposited film of metal shows no dark spots indicating the presence of moisture, a coating of shellac is applied followed by a coating of paint, if desired.

Lead sulphide or galena is a strong metal and adheres tenaciously to the glass. If the mirror shows a grayish color it is usually due to an insufficiently heavy coating of the deposited metal.

An additional coating will remove this defect.

In carrying out the "cold" process, the application of heat is of course omitted and in preparing the final solution no additional distilled water is employed. In other words, the final solution for use in the "cold" process is prepared as follows:

Solution No. 1 1 oz. Solution No. 2 1 oz. Solution No. 3 1 oz.

This final "cold" solution is prepared by adding one part of Solution No. 2 to one part of Solution No. 1. These are thoroughly mixed and allowed to stand for about 15 minutes, after which one part of Solution No. 3 is added. After Solution No. 3 has been added, it is necessary to immediately pour the final solution upon the glass due to the fact that the metal tends to deposit out of solution quite rapidly.

Both the hot and cold processes as hereinbefore described have been found quite effective in the application of a firm and homogeneous film or coating of metallic lead sulphide upon a glass surface or the like, it being of course understood that this lead sulphide is formed by the combination of the sulphur present in Solution No. 3 with the lead present in Solution No. 1. It will be understood that in both the hot and cold processes the thickness of the deposited film or coating may be reduced as desired by introducing additional quantities of distilled water either to the final solution or to the primary solutions.

It is important to note that while galena blue (lead sulphide) will not work or combine with silver it will combine with gold.

Aluminum Mirrors British Patent 433,484

A highly polished aluminum sheet is treated anodically in 2½% borofluoric acid using 20 amp. per sq. ft. at 31 to 33° C., washed and then anodically oxidized in 7% sulphuric acid at 25-26° C. using 12 amp. per sq. ft. After drying, buff with polishing cream.

Silvering Mirrors

a. Silver Nitrate 6 g.
Water 75 cc.
Ammonia (28%) sufficient

Dissolve silver nitrate in water and add sufficient ammonia water to dissolve the precipitate initially formed.

b. Glucose 10 g. Water 100 cc. Mix equal parts of a and b and heat slowly on a steam bath (or in hot water) in the vessel or on the object to be mirrored.

Colored Mirrors

One may use one of two processes to obtain a colored reflecting surface. One process consists of deposition of gold in various thicknesses. The resultant effect of this process is a gold or yellowish to brown colored mirror. This process is limited to a very narrow range of these colors.

A more satisfactory and more widely used process is one where colored glass is used. Pink, red, yellow, purple, green or any desired shade or color glass is used on which silver is precipitated by the regular silvering precipitation process. The silver is then backed on in a normal manner. The resultant effect is a very beautifully colored mirror which is as permanent as the silvering itself. The glass generally used for this purpose is

imported.

Of course, one could use a modification of this colored glass process by spraying or brushing on to the front surface of clear glass a colored transparent coating made up of gum sandarac or similar resin in alcohol and dyed to the proper shade. The back of the glass is then silvered in the normal orthodox method. This type of colored mirror is limited in its life by the durability of the front finish coat. It is also very difficult to obtain a uniform smooth reflecting surface by painting or spraying a finish for during the drying period an orange peel effect may manifest itself on the surface and a wavy condition result.

Matte Silver Finish on Watch Dials Formula No. 1

First clean the article well of oil, grease, etc. Then dip into the following solution:

Sodium Dichromate 4 oz. Concentrated Sulphuric Acid 12 oz. Water 1 gal.

The time of dipping depends on the appearance ultimately desired and must therefore be determined by experiment. Rinse well in water, and silver plate in following:

Silver Cyanide 3.5 oz. troy Sodium Cyanide 4 oz. avoir. Sodium Carbonate

at least 6 oz. avoir.
Water 1 gal.

Finally soak in boiling water to give dead white color.

No. 2

Precipitated silver is used on some types of high grade watch dials where a dead white matte finish is desired. A raised grain effect is obtained at the same time. The following formula can be employed, using precipitated silver:

Precipitated Silver 1 oz. Cream of Tartar 2 oz. Sodium Chloride 2 oz.

Mix dry, add enough water to make thick paste. Apply by running with stiff brush. The proportions may be varied depending upon grain and matte desired. The best results are obtained on alloys rich in copper such as gilding metal.

Sulphur Resisting Alloy German Patent 591,641

 Nickel
 44 to 79 lb.

 Chromium
 9 to 31 lb.

 Aluminum
 at least 9 lb.

 Silicon
 at least 2 lb.

And 0-14% of one or more of the following: Iron, Molybdenum, Copper, Manganese, Carbon.

Alloys for "Tin" Buttons

Lead	16	g.
Antimony	16	
Tin		ø.

Electrical Resistance Wire Alloy U. S. Patent 1,926,213

Gold 58.4 oz. Nickel 41.6 oz.

Heat Treatment of Aluminum Magnesium Silicon Alloy

Anneal for 1 to 3 hours at 500-550° C.; quench in oil or water and temper at 180-250° C. for 1½ to 3 hours.

Corrosion and Heat Resisting Alloy 35% nickel, 15% chromium (balance iron). This material shows very good resistance to oxidation and corrosion at temperatures up to 2000° F., and still retains an appreciable amount of strength.

Improving Babbitt Metal

Babbitt flow characteristics are greatly improved by adding a small amount of rosin to the molten mass.

Zinc Die Casting Alloys

The following zinc die casting alloys are characterized by low metal cost, ease of casting, excellent finish, good resistance to corrosion, permanence of dimensions, and high strength. The percentage limits apply to die castings. Ingot specifications should be narrower.

U. S. Patent 1,596,761

Zamak-2—A.S.T.M. Alloy XXI—S.A.E. Alloy 921

(The name Zamak is trade marked.)
Aluminum 3.5 -4.5%

Grade — 99.99%

Pure) remainder

This alloy is outstanding in hardness, tensile strength, and resistance to corrosion under severe atmospheric exposure conditions.

U. S. Patent 1,779,525

Zamak-3—A.S.T.M. Alloy XXIII— S.A.E. Alloy 903

 Aluminum
 3.5 -4.3 %

 Copper
 0.1 % maximum

 Magnesium
 0.03-0.08%

 Iron
 0.1 % maximum

 Lead
 0.007% maximum

 Cadmium
 0.005% maximum

 Tin
 0.005% maximum

 Zinc (Special High

Grade — 99.99%

Pure) remainder

This alloy is distinguished by excellent retention of impact strength and dimensions.

U. S. Patent 1,852,441

Zamak-5

 Aluminum
 3.5 - 4.5 %

 Copper
 0.75-1.25%

 Magnesium
 0.02-0.08%

 Iron
 0.1 % maximum

 Lead
 0.007 % maximum

 Cadmium
 0.005 % maximum

 Tin
 0.0015% maximum

 Zinc (Special High

Grade — 99.99%

Pure) remainder

The characteristics of this alloy are excellent resistance to corrosion combined with nearly as high strength as Zamak-2 and retention of dimensions nearly equal to Zamak-3.

U. S. Patent Re 18,600

Zamak-6

 Aluminum
 3.5-4.5%

 Copper
 1.0-1.5%

 Magnesium
 0.01 % maximum

 Iron
 0.1 % maximum

 Lead
 0.007% maximum

 Cadmium
 0.005% maximum

 Tin
 0.005% maximum

Grade — 99.99%

Pure) remainder

This alloy offers maximum ease of casting at the expense of maximum resistance to intercrystalline oxidation.

Zinc Slush Casting Alloys

The zinc slush casting alloys offer a desirable combination of high strength, good casting finish, ease of application of plated and other finishes with low metal cost.

Formula No. 1

Zinc (Special High Grade—99.99% Pure).

This metal offers ease of casting and good permanence but lower strength than Formulas No. 2 and No. 3.

No. 2

Aluminum 5-6% Zinc (Special High Grade—99.99% Pure) remainder

This alloy offers the greatest ease of casting and high initial strength but poor permanence.

No. 3

U. S. Patent Re 18,600

Aluminum 4.55-4.95% Copper 0.65-0.85% Zinc (Special High Grade --99.99% Pure) remainder

This alloy is somewhat hard to cast but has good retention of physical properties and high strength.

No. 4

U. S. Patent 1,596,761

 Aluminum
 5.5 -6.5%

 Copper
 2.5 -3.5%

 Magnesium
 0.02-0.1%

 Iron
 0.1 % maximum

 Lead
 0.007% maximum

 Cadmium
 0.005% maximum

 Tin
 0.005% maximum

 Zinc (Special High

Grade - 99.99%

Pure) remainder

This alloy offers the highest strength

and permanence of the zinc base slush casting alloys but is also the most difficult to cast.

Zinc Alloy Solders Formula No. 1

Cadmium		82.5%
Zinc		17.5%
Melting Point		508° F.
This solder is	most	advantageously

used in soldering zinc alloy castings containing aluminum. No flux is necessary.

Note: In making this solder, solid cadmium should be added to molten zinc since cadmium fumes have a very dangerous toxic effect. If the cadmium be melted separately, the temperature should not be allowed to rise above 660–700° F. and the surface of the molten metal should be treated with a flux of ammonium chloride.

U. S. Patent 1,988,010 Percentage by Weight

Composition	Tin	\mathbf{Zinc}	Cadmium	Freezing Point ° F.
Formula No. 1	20	53	27	617
No. 2	20	48	32	604
No. 3	30	53	17	630
No. 4	30	46	24	599
No. 5	30	42	28	595
No. 6	40	36	24	581

The above solders are used principally for soldering aluminum and aluminum base alloys. They may be used with or without fluxes depending on the cleanliness of the metal parts.

Cleaning of Zinc and Zinc Alloys

The successful application of plated and other coatings to zinc, zinc alloy die castings, and zinc alloy slush castings depends largely on the suitability and effectiveness of the method of cleaning used.

Cleaning may be accomplished by any one of three methods: (1) Mechanical cleaning by means of sandblasting or scratch brushing, (2) alkaline cleaning and (3) solvent cleaning.

Mechanical Cleaning

Sandblasting with 80 to 100 mesh abrasive is probably most effective since it simultaneously removes grease and dirt and roughens the surface of the metal.

Alkaline Cleaning

Alkaline cleaning has been accomplished very effectively by the use of trisodium phosphate in concentration of 6 oz. per gal. of water. This solution when used at or near the boiling temperature and with sufficient current from a 6-volt source to cause violent gassing with the work as the cathode, should remove all grease and oil in ½ to 2 minutes. Alternate hot and cold rinses followed by a brief immersion in 10%

hydrochloric acid and a final rinse in hot water to facilitate drying will effectively remove the film of alkaline cleaning salts and present a surface suitable for plating or other finishes.

Soldering Zinc and Zinc Alloy Castings

Zinc may be soldered easily, using ordinary solder and a flux consisting of acidulated zinc chloride or killed muriatic (hydrochloric) acid.

(hydrochloric) acid. Zinc alloys containing aluminum are quite difficult to solder, requiring the use of a solder consisting of the cadmiumzinc eutectic (82.5% cadmium—17.5% zinc—melting point 508° F.).

Machining Zinc and Zinc Alloy Castings Both rolled zinc and zinc alloy castings are machined most advantageously by using tools with more rake than is customary in machining other common metals. The cutting tool should have 15-20° rake and 6-8° clearance.

Two-fluted drills with spiral angles about twice the usual 24 degrees are satisfactory. The included angle of the cutting edges may be advantageously reduced. The clearance angle should be 15 degrees at the periphery of the drill and gradually increased still further as the drill point is approached. Beveling off the end of the flute back of each cutting edge provides more chip clearance for rapid work.

Soapy water is ordinarily a satisfactory lubricant. Kerosene may be used as a lubricant to insure satisfactory separation of thing

tion of chips.

Low Temperature Glaze for Art Ware and Enameled Brick

White Lead	35 lb.
Feldspar	17 lb.
Flint	20 lb.
Whiting	8 lb.
China Clay	8 lb.
Colemanite	12 lb.
Tin Oxide	
	5 lb.
Matte Glaze—Cone 06 to	Cone 02:
White Lead	100 77
	490 lb.
Whiting	490 lb. 138 lb
	138 lb.
Whiting Cornwall Stone	138 lb. 114 lb.
Whiting Cornwall Stone China Clay	138 lb. 114 lb. 210 lb.
Whiting Cornwall Stone	138 lb. 114 lb.

For light green use 2 to 3% copper oxide; for light brown 2% manganese dioxide; for blue 1% cobalt oxide; for yellow 2% sodium uranate; for yellow brown ½ to 2% Crocus Martis.

Atware Satin Glaze-Cone 04:

White Lead	410	lh
Flint	227	
Feldspar	85	
Zinc Öxide	90	
Tin Oxide	60	
Barium Carbonate	42	
Titanium Dioxide	32	
China Clay	54	lb.
Green Matte Glaze-Cone 2:		

Green Matte Glaze—Cone	2:
Red Lead	165 lb.
Feldspar	222 lb.
Whiting	40 lb.
Zinc Oxide	32 lb.
Copper Oxide	12 lb.
Calcined Georgia Kaolin	55 lb.
English Ball Clay	64 lb.

This gives a good wax-like texture green for artware or enameled brick.

Vitreous Enameling Process British Patent 411,380

A mixture of spinel-forming materials, e.g., water 100, ferric oxide 5, nickel oxide 4, calcium fluoride 20, boric acid 45, clay 10 parts, is applied to the iron surface (not necessarily free from rust) and heated at 750-800° for a few minutes in an atmosphere of reduced oxygen content (admixture of producer or waste gases, etc.).

White Vitreous Enamel U. S. Patent 1,933,437

A white enamel for sheet iron and hollow-ware comprises flint 29.236, borax 13.127, sodium nitrate 5.727, sodium carbonate 10.740, red lead 14.920, barium

carbonate 7.757, calcium fluoride 6.563, antimony oxide 4.773, and sodium antimonate 7.160%.

Spark Plugs French Patent 772,601

A ceramic product for spark plugs is composed of a difficultly fusible oxide, e.g., corundum, and a binder which during thermal expansion behaves elastically toward the oxide used. The binder should become plastic at 500-800° C. An example of a binder for use with corundum contains steatite or talc 32.7, kaolin 43.3 and feldspar 24 parts by weight.

Synthetic Precious Stones (Spinels) U. S. Patent 1,952,255

(a) Artificial alexandrite is made by fusing aluminum oxide 85 and magnesium oxide 15% containing cobalt 0.06, iron 0.04%, and vanadium 0.04%, and (b) a violet spinel by fusing the same aluminum oxide-magnesium oxide mixture with iron 1.5 and cobalt 0.005%.

Corundum Abrasive Crystals U. S. Patent 1,966,406

A mixture of raw materials is prepared consisting of aluminous ore such as bauxite or diaspore, silica sand, and an addition agent such as magnesia so proportioned as to give the following ratio of important ingredients:

 Alumina
 70 lb.

 Silica
 25 lb.

 Magnesia
 5 lb.

This mixture may be fused in an electric furnace of the steel shell are type commonly used in the artificial abrasive industry. The ratio of power input to application of the mix is observed closely as means of governing the temperature of the melt. Thus, under any given rate of power input, a fast feed produces a relatively cool meit, whereas a retarded feed tends to produce a relatively hot The temperature of the melt at the time of withdrawal of the power determines the size and distribution of the corundum crystals. The cool melt produces small crystals uniformly spread through the matrix whereas the hot melt gives rise to the development of large crystals, in pocket formation in the mass.

After the shell has been charged to its capacity and the fusion is completed the electrodes are withdrawn and the cooling process allowed to proceed normally.

Brick Glazing

White	Enamel	Batch	Weights
ed Lead			12
*** * . *			

Red Lead	125.4
Whiting	3 5
No. 419 Feldspar	66.1
Raw Kaolin	12.9
Calcined Kaolin	6.7
Flint	40
Tin Oxide	30

Black Enamel

To the above base enamel batch, without the tin oxide, the following is added:

Cobalt Oxide (CoO)	6
Iron Oxide (Fe ₂ O ₃)	8
Manganese Dioxide (MnO ₂)	2

Blue Enamel

Batch weights of base enamel:

Buckingham Spar	66.32
Red Lead	120.84
Whiting	36
Tin Oxide	57.77
Raw Clay (Kaolin)	12.9
French Flint	40.34
Calcined Kaolin	11.22
To the above base is added:	
Black Oxide of Copper	12
Black Oxide of Cobalt	18
Black Oxide of Nickel	6

Brown Enamel

To the above base is added:

Red Oxide of Iron (Fe₂O₃) 16

The production of other colors is merely a matter of experiment with the addition of coloring oxides.

These glazes contain tin oxide as an opacifier and on a smooth body develop a glossy enamel of sufficient weight to perfectly mask the red of the shale brick.

Slips

95% Tennessee Ball Clay (for white slip use English Ball Clay).

5% Sodium Chloride are of simple

materials and easily made up.
For green slip add 20% Chromium
Oxide (Cr₂O₃) to the above base. The
batch then is:

Cone 02 to Cone 2

Clay	3 3 3 3 3	380
Sodium Chloride		20
Chromium Oxide		80

For blue slip, add 6% Cobalt Oxide to base. Batch:

Clay	380
Sodium Chloride	20
Cobalt Oxide	24

Stone Waterproofing

An economical treatment that is very durable may be made by dissolving from 6 to 12 oz. of a high-melting-point paraffin to the gallon of solvent, such as mineral spirits, naphtha, gasoline, etc. This usually gives high waterproofing values on materials of medium to coarse textures. For fine-pore structures it will be desirable to add from 3 to 6 oz. of china wood oil to the gallon of gasoline.

Stucco Waterproofing U. S. Patent 1,942,601

Sodium Stearate 5 lb. Water 95 lb. Warm to 50° C. and stir till uniform,

then add
Suet 2 lb.
Cresol Emulsion 4/4 oz.

Masonry Waterproofing British Patent 413,463

Spermaceti	4 lb.
Paraffin Wax	1 lb.
Rubber	1 lb.
Mineral Spirits	25-50 lb.
Trichloroethylene	25-50 lb.
Stir until dissolved.	

Vitreous Slips for Brick, Terra Cotta and Roofing Tile

Buff	
Fireclay	130 lb.
Shale	100 lb.
White Lead	40 lb.
Blue	
Ball Clay	200 lb.
Cobalt Oxide	9 lb.
Manganese Dioxide	6 lb.
White Lead	50 lb.
Green	
Ball Clay	200 lb.
White Lead	50 lb.
Chrome Oxide	40 lb.
Manganese Dioxide	24 lb.
Cobalt Oxide	5 lb.
Black	
Ball Clay	60 lb.
Blackbird Clay	140 lb.
White Lead	30 lb

Mix the above materials with sufficient water to make a heavy slip, and apply either by spraying or brushing on the dry body, then fire.

CERAMIC RAW MATERIALS

Chemical Constants

Per Cent Smelt Loss	34.6	22.3		25.5	44	14	68.4 74.3 73.3	35.5		71.6	
Melting Point Deg. C.	D770 red heat	D900, M1360 820–860 550	732	577 above 1426	D825	O O	80.75 56 96.8	1235			
Veight	156 342.1 291.5	197.4 466 339.7	190.6	69.6 172.4 128.4	144.4 100.1	128.6 237.8	298 582.2 562.2	124 79.6	92.9	/8.1 	323
Equivalent Weight		RO_2	$^{ m R}_{ m 2}{ m O}_{ m 3}$			$^{\rm RO_2}_{\rm R_2O_3}$	$\substack{R_2O_3\\R_2O_3}$		${ m RO}_2^2$		
	R203	$ \begin{array}{c} RO & 197.4 \\ RO & 197.4 \\ RO & 3 \\ RO & 4 \\ RO & 3 \\ $	RO 381.2 RO 201.3	K2O3	RO	$^{ m R}_{ m 2}{ m O}_3$ 258.1 $^{ m R}{ m O}$ 118.9	RO 291.1 RO 281.1		8 R ₂ O ₃ 556.8 5 R ₂ O ₃ 524.5	PEO	RO.5R ₂ O ₃ 2RO. R ₂ O ₃
Molecular Weight	156 342.1 291.5 197.8	197.4 466 339.7 310	340 381.2 201.3	69.6 172.4 128.4	144.4 100.1 152	258.1 118.9	291.1 281.1 75	124 79.6	556.8 RO 556.8 524.5 RO 524.5	78.1 159.7 278	323
Formula		$egin{array}{l} \operatorname{BaCO}_3 \ \operatorname{Bi}_2 O_3 \ \operatorname{SD}_2 S_3 \ \operatorname{Ca}_3 \left(\operatorname{PO}_4\right)_3 \end{array}$	ိုင			O _s	$\begin{array}{c} {\rm Co(NO_3)_2.6H_2O} \\ {\rm CoSO_4.7H_2O} \\ {\rm CoSO_4.7H_2O} \\ {\rm CoOO} \end{array}$		\sum_{2}^{2}	$^{\mathrm{Ca.r.2}}_{\mathrm{Fe2O_3}}$ $^{\mathrm{FeSO_4}}_{\mathrm{4.7H_2O}}$	
Material	Aluminum Hydroxide Aluminum Sulphate Ahtimony Oxide Arsenic Oxide	Barium Carbonate Bismuth Oxide Black Needle Antimony Bone Ash	Bone Ash Borax Borax (Melted)	Cadmium Carbonate	Cadmum Surpinde Calcium Carbonate Chromium Oxide	Clay (China)	Cobalt Nitrate Cobalt Sulphate Cobaltcons Oxide	Copper Carbonate Copper Oxide (Cupric) Copper Sulphate	Feldspar (Potassium) Feldspar (Sodium)	Iron Oxide (Ferric Oxide)	Lead Chromate

$egin{array}{c} -rac{59.4}{18.4} \ m{H_2O} = 36.4 \ m{52.1} \end{array}$	$\frac{83.8}{H_2O = 44.9}$	31.8 16.3 53.4 2.3	24.6	12.1 63.5 $$ 12.1 $$ $$	29.8	81.4 123 2700 183.3 2550 Courtsey of Eureka Plint and Spar Co., Inc.
618–710 O ₂ 535 650 D350	2800 O ₂ 400	896 397.5 360.4 337 D500	688 1600–1750 ———	310	D1075 1127 1560 D	2700 2550 f Eureka Flint c
73.9 86.9 197.9 84.3	40.3 246.5 280.9 74.7	138 147.1 112.2 202.2 228.5	79.2 60.1 511.6 286	298 85 188 322.2	147.6 94.8 150.7 80.1 258.5	81.4 123 183.3 ourtsey o
		RO_3			RO_2	,0
RO RO RO RO	RO RO	RO 294.2 R_2O_3 294.2 RO RO RO RO	RO ₂		RO 126.4 RO 126.4 RO ₂ RO ₂	$^{ m RO}_2$
223.2 73.9 86.9 197.9 84.3	40.3 246.5 280.9 74.7	138 294.2 56.1 101.1 685.6	633.6 60.1 511.6 286	298 298 322.2	348.2 147.6 379.3 150.7 80.1	81.4 123 183.3
$\begin{array}{c} \text{PbO} \\ \text{Li}_2\text{CO}_3 \\ \text{MnO}_2 \\ \text{MnCl}_2 \cdot 4\text{H}_2\text{O} \\ \text{MgCO}_3 \end{array}$	$egin{array}{l} m MgSO_4\cdot 7H_2O \\ m NiSO_4\cdot 7H_2O \\ m NiO \\ m NiO \end{array}$	$egin{array}{c} K_2CO_3 \ K_2Ch_2O_7 \ KOH \ KNO_3 \ Pb_3O_4 \end{array}$	$\frac{\text{Se}_{8}}{\text{SiO}_{2}}$ $\frac{\text{SiO}_{2}}{2\text{NaSbO}_{3}.7\text{H}_{2}\text{O}}$ $\frac{10\text{H}_{2}\text{CO}_{3}.10\text{H}_{2}\text{O}}{2\text{Na}_{2}\text{CO}_{3}.10\text{H}_{2}\text{O}}$	$ m Na_{2}Cr_{2}O_{7} = Na_{2}CO_{7} = Na_{2}O_{7} = Na_{2}O_{4} = Na_{2}SO_{4} $	${ m Na_2UO_4} \\ { m Na_2UO_4} \\ { m SrCO_3} \\ { m 3MgO-4SiO_2\cdot H_2O} \\ { m SnO_2} \\ { m TiO_2} \\ { m 2PbCO_3\cdot Pb(OH)_2} \\ { m 2PbCO_3\cdot Pb(OH)_2} \\ { m Na_2UO_3\cdot Pb(OH)_2} \\$	$Z_{\rm nO}$ $Z_{\rm rO_2}$ $Z_{\rm rSiO_4}$
Litharge Lithium Carbonate Manganese Dioxide Manganous Chloride Magnesium Carbonate (Magnesite).		Potassium Carbonate Potassium Dichromate Potassium Hydroxide Potassium Nitrate Red Lend	Selenium Silica (Flint) Sodium Antimonate Sodium Carbonate (Crystal)	Sodium Carbonate (Fused) Sodium Dichromate Sodium Nitrate Sodium Silico Fluoride Sodium Sulphate	Sodium Uranate	Zinc Oxide Zirconium Oxide Zirconium Silicate

FUSING TEMPERATURES OF CERAMIC RAW MATERIALS

Material	Formula	Temperature Deg. C.
Aluminum Oxide (Alumina) Antimony Oxide Arsenic Oxide Barium Oxide Barium Oxide Bone Ash Borax (Melted) Boric Acid Boric Oxide Calcium Fluoride (Fluorspar) Calcium Oxide (Lime) Calcium Phosphate Calcium Silicate (Wollastonite) Cerium Oxide Chromium Oxide Cobaltous Oxide Copper Oxide (Cupric) Feldspar, Potassium Feldspar, Sodium Fluorspar Iron Oxide (Ferric Oxide) Kryolith Lead Oxide Lead Silicate Lithium Oxide Magnesium Oxide Magnesium Oxide Manganous Oxide Nickelous Oxide Phosphoric Oxide Potassium Silicate Silica (Flint)	Al ₂ O ₃ Sb ₂ O ₃ As ₂ O ₅ BaO 4Ca ₃ (PO ₄) ₂ ·CaCO ₃ Na ₂ B ₄ O ₇ B ₂ O ₃ B ₂ O ₃ CaF ₂ CaO Ca ₃ (PO ₄) ₂ CaSiO ₃ CeO ₂ Cr ₂ O ₃ CoO CuO K ₂ O·Al ₂ O ₃ ·6SiO ₂ Na ₂ O·Al ₂ O ₃ ·6SiO ₂ CaF ₂ Fe ₂ O ₃ Na ₃ AlF ₆ PbO PbO·SiO ₂ Li ₂ O MgO MnSiO ₃ MnO NiO P ₂ O ₅ K ₂ O·SiO ₂ SiO ₂	Deg. C. 2050 1550 200 02450 732 577 577 1300 2570 1550 1540 1950 196 1235 1170-1235 1120-1215 1300 1565 888 766 2800 1273 1650 02400 563 red heat 976 1710
Potassium Silicate	$K_2^{2}O \cdot SiO_2$	976
Soda Ash (Sodium Carbonate) Sodium Antimonate	Na ₂ CO ₃ 2NaSbÖ ₃ ·7H ₂ O	851
Sodium Oxide	$egin{array}{l} { m Na_2SiO_3} \end{array}$	$rac{ ext{red heat}}{1080}$
Sodium Silico Fluoride Tin Oxide (Stannic)	Na_2SiF_6 SnO_2	1127
Titanium Oxide Zinc Oxide Zirconium Oxide	$egin{array}{l} ext{TiO}_2 \ ext{ZnO} \ ext{ZrO}_2 \end{array}$	1560 1800 2700
	Country of Theorem	

Courtsey of Eureka Flint and Spar Co., Inc.

Cold Tile and Brick Glaze U. S. Patent 2,019,980

Portland Cement 10 parts by vol. Iron Oxide 1 part by vol. Calcium Stearate and Water (1-2%) 5 parts by vol.

Mix thoroughly and pass through a

screen to remove lumps.

The glaze is now ready for application to the product or article, which, for example, may be cement tile, building blocks, or other suitable materials. This may be accomplished by brushing, dipping or spraying the glaze thereon until the desired coating is effected. The

glazed objects may be trimmed and then placed in a curing chamber which is kept moist for several days. In order to get best results, the tiles are thereafter placed in storage for a week or longer, to age or cure, until the permanent hardening or setting of the glaze is completed.

Enamel Ware Undercoat U. S. Patent 1,962,617

The base metal is sprayed with a suspension of

Cobalt Oxide		3	oz.
Bentonite		1.5	oz.
Water		100	oz.

CUBICAL COEFFICIENTS OF EXPANSION OF CERAMIC RAW MATERIALS

${f Material}$	Formula	2	X 10-7
Aluminum Oxide (Alumina)	Al_9O_3	(0.52)	5.0
Antimony Oxide	Sb_2O_3	, ,	3.6
Arsenic Oxide	As_2O_5		2.0
Barium Oxide	BaŌ	(5.2)	3.0
Bone Ash	$4\text{Ca}_3(\text{PO}_4)_2 \cdot \text{CaCO}_3$	` ,	-
Borax (Melted)	Na ₉ B ₄ O ₇		3.16
Boric Acid	$\mathrm{B}_2\mathrm{\bar{O}}_3$	(-1.98)	0.1
Calcium Fluoride (Fluorspar)	CaF ₂	` ,	2.5
Calcium Oxide (Lime)	CaO		5.0
Calcium Phosphate	$Ca_3(PO_4)_2$		3.65
Cerium Oxide	CeO_2		4.2
Chromium Oxide	$Cr_2\bar{O_3}$		5.1
Cobaltous Oxide	C_0O		4.4
Copper Oxide (Cupric)	CuO		2.2
Fluorspar	CaF_2		2.5
Iron Oxide (Ferric Oxide)	$\mathrm{Fe_2O_3}$		4.0
Kryolith	Na ₃ AlF ₆		7.4
Lead Oxide	PbO	(3.0)	4.2
Lithium Oxide	$\mathrm{Li_2O}$		2.0
Magnesium Oxide	$\overline{\text{MgO}}$	(1.35)	0.1
Manganous Oxide	MnO		2.2
Nickelous Oxide	NiO		4.0
Phosphoric Oxide	P_2O_5		2.0
Potassium Oxide	$K_2^{\circ}O$	(11.7)	8.5
Silica (Flint)	SiO_2	(0.15)	0.8
Sodium Antimonate	$2 \text{NaSbO}_3 \cdot 7 \text{H}_2 \text{O}$		
Sodium Oxide	Na_2O	(12.96)	10.0
Sodium Silicate	$\mathrm{Na_{2}SiO_{3}}$		2.96
Sodium Silico Fluoride	Na_2SiF_6		5.0
Tin Oxide (Stannic)	SnO_2		2.0
Titanium Oxide	${ m TiO_2}$		4.1
Zinc Oxide	ZnO		1.8
Zirconium Oxide	${ m ZrO_2}$		2.1

Courtsey of Eureka Flint and Spar Co., Inc.

Pottery Glaze French Patent 44.786

TIGHON TAVONO 1191	00
Feldspar	26 kg.
Quartz	2 kg.
Minium	49 kg.
Barium Borosilicate	15 kg.

This is applied with coloring materials after grinding, without fritting.

Flooring Tile Norwegian Patent 55,221

The mass before drying is composed of linseed oil 7, coal tar 1, alkali silicate 1, varnish 1, water 1, glue 1, cement 1, quartz sand 5, clay 1 and salt 1 part. all by weight.

Colored Roofing Granules U. S. Patent 1,944,294

Burned clay granules are impregnated with arsenic trioxide and surface washed. Then treat with 15% basic copper acetate solution, wash and dry.

Manufacture of Light-Weight Ceramic Tile

U. S. Patent 1,925,985

A mixture of ball clay 45-65 (56.7), plaster of Paris 10-20 (13.1), and sawdust (I) 25-40 (32.1) is rendered plastic by addition of 80-120 (103)% of water and cast into waxed molds. The dried tiles are heated for 4 hours at about 500° F. until (I) is charred, then slowly (4 hours) up to 1200°, at which temperature they are kept for 4 hours until the carbon is burnt out and shrinkage ceases.

White Enamel for Wire U. S. Patent 1,938,691

Fuse together:				
Borax		16.5	lb.	
Feldspar	50-	-45	lb.	
Silica		9.2	lb.	
Soda Ash	20-	-25	lb.	
Sodium Nitrate		3	lb.	
			-	

Quench and grind with 8% titanium dioxide.

Light Weight Refractory U. S. Patent 1,945,232

Brick or Pottery Clay 1 lb.
Rice Hull Ashes 2 lb.
When the above is fired the product is 40% lighter than usual.

Synthetic Lumber U. S. Patent 1,974,277

Magnesium Oxide	30 1ե.
Aluminum Oxide	20 lb.
Sawdust	50 lb.
Beach Sand	10 lb.

(Continued on page 245,

TEMPERATURE EQUIVALENT OR FUSING POINTS OF CONES (Adopted from Table XIII, of U. S. Bureau of Standards' Report)

Note: These approximate values are given by the Bureau of Standards to the nearest 5° C. from the average determinations.

accorded to the accorded to		02254			
Soft Series:	****				
~	When Fired Slowly			When Fired Rapidly	
Cone Number	20° C. 1	er Hour		150° C. p	
	° Cent.	° Fahr.		° Cent.	° Fahr.
022	585	1085		605	1121
021	595	1103		615	1139
020	625	1157		650	1202
019	630	1166		660	1220
018	670	1238		720	
					1328
0 7 .3	720	1328		770	1418
152	735	1355		795	1463
015	770	1418		805	1481
014	795	1463		830	1526
013	825	1517		860	1580
$012 \dots \dots \dots$	840	1544		875	1607
011	875	1607		905	1661
Low Temperature Series:					
010	890	1634		00~	1040
09	930			895	1643
0.0		1706		930	1706
	945	1733		950	1742
07	975	1787		990	1814
06	1005	1841		1015	1859
05	1030	1886		1040	1904
04	1050	1922		1060	1940
03	1080	1976		1115	2039
02	1095	2003		1125	2057
01	1110	2030		1145	2093
Intermediate Temperature Series	:				
1	1125	2057		77.00	0100
	1135	2075		1160	2120
				1165	2129
	1145	2093		1170	2138
4	1165	2129		1190	2174
5	1180	2156		1205	2201
<u>6</u>	1190	2174		1230	2246
7	1210	2210		1250	2282
8	1225	2237		1260	2300
9	1250	2282		1285	2345
10	1260	2300		1305	2381
11	1285	2345		1325	2417
12	1310	2390		1335	2435
13	1350	2462		1350	2462
14	1390	2534		1400	2552
$1\overline{5}$	1410	2570		1435	
16	1450	2642			2615
12 2				1465	2669
	1465	2669		1475	2687
# A	1485	2705		1490	2714
00	1515	2759		1520	2768
20	1520	2768		1530	2786

TEMPERATURE EQUIVALENT OR FUSING POINTS OF CONES—Continued

	High	Temperature	Series:
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	When	Heated	at 100° per Hour
		° Cent.	° Fahr.
23	 	1580	2876
26	 • • •	1595	2903
27	 	1605	2921
28	 • • •	1615	2939
29	 	1640	2984
30	 	1650	3002
31	 	1680	3056
32	 	1700	3092
33	 	1745	3173
34	 • • •	1760	3200
35	 • • •	1785	3245
36	 	1810	3290
37	 	1820	3308
38	 	1835	333 5
*39	 • • •	1865	3389
40	 • • •	1885	3425
41	 • • •	1970	3578
42	 • • •	2015	3659

^{*} The last four cones were heated at 600° per hour.

Moisten with magnesium chloride solution and calcium magnesium chloride and after forming dip in a solution of magnesium silicofluoride and potassium sulphate.

Building Material Austrian Patent 137,323

A fibrous organic material 3, a pulverulent mineral 5-7.5 and water-glass solution of 36-38° Bé. 5-7.5 parts are mixed together, molded in a perforated mold, and dried. The organic material may be wood pulp, straw or sugar-cane waste and the mineral may be asbestos or kaolin.

Composition for Floors and Wall Surfaces

Austrian Patent 137,328

Dried sawdust 40-60, cement 30-40 and lime 5-10 parts are kneaded with 50-70 parts of concentrated water-glass solution. The dried composition can be subjected to the same mechanical treatments as wood.

Artificial Gypsite Plaster U. S. Patent 1,932,120

Gypsum	26,180 lb.
Dry Peat	600 lb.
Clay	2,820 lb.

Stir and heat with calcium chloride solution (d. 1.4) 4 qt. in a plaster kettle heating at 155-165° C.

Courtsey of Eureka Flint and Spar Co., Inc.

Opal Vitreous Marble, Artificial French Patent 784,067

Sand	500	kg.
Soda Ash	200	kg.
Lime	100	kg.
Sodium Nitrate	20	kg.
Spar	60	kg.
Feldspar	70	kg.
Antimony	1.6	kg.
Arsenic	0.2	kg.
Manganese	0.6	kg.
Zinc Oxide	20	kg.
Fish Offal or Blood	200	kg.

Artificial Marble Formula No. 1

British Patent 416,774

White Cement		50	lb.
Marble Dust		50	lb.
Calcium Carbonate	Powder	3.85	lb.
Calcium Oxalate		0.50	lb.
Borax		0.15	lb.
Starch		0.60	lb.

Mix with water and allow to set.

No. 2

British Patent 430,948

Magnesium Oxide	100 lb.
Marble Dust	30 lb.
Calcium Sulphate Dust	20 lb.
Make into a paste with	magnesiun
ploride (d. 120-126) and	then add

Magnesium Oleate		1	lb.
Magnesium Stearate	28 MARS	1	lb.
Tallow Soap Solution	1 (2%)	10	lb.

STANDARD SCALES FOR TESTING SIEVES

		Di- ameter of Wire (Inches)	.148	.135	105	.105	.092	.088	.065	.065	.044	.0328	.032	.033	.035	.028 700	.020	.0172	.0141	.0125	0118	0100	20092	
es		Mesh (Per Lineal Inch)		1		ļ	1	27. %	$3\frac{1}{2}$	4	ທ	-	œ	6	01	77	14	20	24	828	90 e 91 fd	94 95	84 65)
Sieve Seri		(Fractions of Inch) Approx.)	1 7/8	% ⁷	7% 17%	2/ ₁ 6	%	5/16 1/2	7/4 289	$\frac{3}{16}$	5/35 1/3	8 ½	**************************************	204 704	$\frac{1}{16}$	1 8	\$	1/39	-	1	>	⁷ 07		
Tyler Standard Sieve Series		Openings (Milli- meters) (26.67 22.43	18.85	13.33	11.20	9.423	7.925	5.613	4.699	3.962	2.794	2.362	1.981	1.651	1.397	1.168 991	.833	.701	.589	.495	351	295	1
Tyler 8	For Closer Sizing	Sieves from .0029 to 1.050-in. Ratio $4\sqrt{2}$ or 1.189	1.050	.742	.624 525	.441	.371	.312	221	.185	.156	110	.093	.078	.065	.055	.046	.0328	.0276	.0232	.0195	0138	.0116	1
	Tyler Standard	$\sqrt{2}$ or 1.414 (Openings in Inches)	1.0	.742	595	1	.371	696	9	.185	161	Ter.	.093	-	.065	13	.046	.0328	1	.0232	5	*0TO	.0116	1
		Wire Diameter (Milli- meters)	1.85	1.45	1.27	1.02	.92	.84 27	5.69	.61	54.	6.4	.37	.53	.29	.25	22.	162	.140	.119	707	074	.063	
	eries	Wire Diameter Diameter (Milli-	.073 1.85 .065 1.65																					
	d Sieve Series	_ H		.057	.050	.040	.036	.0331	.0272	.0240	.0213	0165	.0146	.0130	.0114	8600.	.0087	.0064	.0055	.0047	.0040	0003	.0025	
	S. Standard Sieve Series	Wire I Diameter (Inches)	.065	5.66 .057	4.76 .050	3.36 .040	2.83 .036	2.38 .0331	1.68 .0272	1,41 .0240	1.19 .0213	84 .0165	.71 .0146	.59 .0130	.50 .0114	.42	.35 .0087	.250 .0064	.210 .0055	.177	. 149 .0040	105 0009	.088 .0025	
	U. S. Standard Sieve Series	Sieve Opening Wire I (Milli- Diameter meters) (Inches)	8.00 .073 6.73 .065	.223 5.66 .057	.187 4.76 .050 157 4.00 044	132 3.36 .040	.111 2.83 .036	.0937 2.38 .0331	.0661 1.68 .0272	.0555 1.41 .0240	.0469 1.19 .0213	.0594 1.00 .0169 0331 .84 .0165	.0280 .71 .0146	.0232 .59 .0130	.0197 .014	.0165 .42 .0098	7800. 35 .0087	.0098 .250 .0064	.0083 .210 .0055	.0070 .177 .0047	.0059 .149 .0040	0049 .125 .0054	.0035 .088 .0025 0029 074 0021	
	က်	Sieve Sieve Opening Wire I Opening (Milli- Diameter r (Inches) meters)	.315 8.00 .073 .265 6.73	34, .223 5.66 .057	4 .187 4.76 .050 5 157 4.00 044	6 .132 3.36 .040	7 .111 2.83 .036	8 .0937 2.38 .0331	12 .0661 1.68 .0272	14 .0555 1,41 .0240	16 .0469 1.19 .0213	18 .0394 1.00 .0139 20 0331 84 0165	25 .0280 .71 .0146	30 .0232 .59 .0130	35 .0197 .50	40 .0165 .42 .0098	50 .0138 .35 .0087	600. 0098	70 .0083 .210 .0055	80 .0070 .177 .0047	100 .0059 .149 .0040	120 .0049 .125 .0054 140 .0041 .105 .009	170 .0035 .088 .0025 200 0029 074 0021	

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.0056	8800	.0026	.0024	.0021	ት በ	.807	.192	.148	.0029 inch he standard s, the open-1.414. s the Tyler this series it his series in root of 2 Spar Co., Inc.
100	115	150	170	200	coarser sizing-3- to 11/2-inch opening	an demonstrated in	-	-	This sieve scale has as its base an opening of .0029 inch which is the opening in 200-mesh .0021-inch wire, the standard sieve, as adopted by the U. S. Bureau of Standards, the openings increasing in the ratio of the square root of 2 or 1.414. Where a closer sizing is required column 2 shows the Tyler Standard Screen Scale with intermediate sieves. In this series the sieve openings increase in the ratio of the fourth root of 2 or 1.189.
11	1	1	1	1	- to 11/2	ഹ	c 3	$1\frac{7}{2}$	base an sh0021 Bureau te square uired co ermediat he ratio
.175	.124	.104	.088	.074	sizing—3	1		l	s as its 1 200-me te U. S. tho of th ng is red with int case in tl
.0069	.0049	.0041	.0035	.0029	For coarser		I	i	e scale has opening in pted by the refuser sizir closer sizir reen Scale mings incre
.0058		.0041	I	.0029	F	1		1	This sieve which is the sieve, as adoings increasing. Where a Standard Scithe sieve ope or 1.189.

of 2, or 1.1892 as the ratio of the width of one opening to the next smaller opening. In making selections from this series it is recommended that this be done on some systematic plan, as for below this in the series are related to it by using the fourth root example, the selection of every other sieve or of every fourth one

the sieves above and

mm. is the basic one, and essentially metric.

scale is

The sieve having

Stone Wood Composition Flooring British Patent 426,739

A 1:9 mixture of sodium thiosulphite and calcium carbonate is added to plaster of paris containing sawdust or cork filler. Proportions by volume of 1:5:4 respectively are preferred.

Artificial Stone Flooring U. S. Patent 1,968,784

Calcined Ma	gnesite	100	lb.
Sawdust		200	lb.
Fine Stone	Screenings	50	lb.
Magnesium	Chloride	100	lb.
Emulsified A	Asphalt	$1\frac{1}{2}$	gal.
Water	to make a	soft m	ortar

"Eternit" Artificial Slate

Cement	100 lb.
Asbestos	20–25 lb.
Pigment	5 lb.
Rosin Solution	to suit

Artificial Stone British Patent 430,404

(a) Portland cement 2 and clay dust 0.25 are mixed, (b) cement 2 and slate dust 1 are mixed therewith, (c) cement 2 and shale 1 are mixed therewith, (d) cement 2 and river of sea sand 1 parts are mixed therewith, (e) the calcium oxide solution is added, 0.5 gal. at a time, with mixing after each addition. (7 lb. of calcium oxide and 2 gal. of water may be used for each 28 lb. of cement), (f) the product is molded into slabs, dried 24 hours and then baked 2 hours at 100° C. The slabs are then painted, heated 1 hour, cooled 30 minutes, coated with enamel or cellulose paint, heated 2 hours at 150°, smoothed with cuttlefish bone and polished.

Wall Board, Artificial U. S. Patent 1,976,190

Calcined magnesite 12 parts; sawdust 3 parts; an aqueous solution of magnesium chloride at about 18° Bé., 14 parts; molasses 4 of 1 part; said ingredients being combined in a creamy fluid mixture sufficiently thin to be readily poured into a form or mold.

Commercial Porcelain

The clay used in making porcelain varies in each locality and it will thus be necessary in the following formulas to include the chemical analysis of the clay

and spar used. The physical properties of various mixtures are best illustrated by the triangular diagram shown by Gilchrest and Klinefelder in the Electric Journal, March and April, 1918. This diagram shows the variation of mechanical, electrical and thermal properties with variation in mixture.

The usual mixtures for electrical porcelain is 40-50% clay, 25-30% spar and 25% quartz. The European insulator materials are ground extremely fine and fired to a hard glass-like body, usually

Seger cone 14-16.

German Porcelain Mixtures Insulator Porcelain

Ka	olin	fro	m Ha	lle	50.	4%
Cla	уf	rom	Halle	3	31.	5%
Spa	ar				18.	1%
Ea	Iren	ŧο	530%	clav	substance	290

quartz and 18% spar.

Household Porcelain

Formula.	No. 1	No. 2	No. 3
Kaolin from Halle	55%	60 %	22.4%
Clay from Halle	27%		44.8%
Zettlitzer Kaolin		18 %	16.2%
This equals			
Clay Substance	54%	55 %	60 %
Quartz		22.5%	22.5%
Spar	18%	22.5%	17.5%

The kaolin from "Halle" mines contains about 61.77% clay substance, 37.84% quartz and .39% spar. The clay from the same mines are about 70% clay substance and 30% quartz.

Karlsbaden Czechoslovakian Porcelain Clay Substance Quartz 29.62%-24.5 % 17.26%-21.93% Spar Calcium Carbonate 1.25%- 1.6 %

Danish Porcelain

Clay Substance	31.8%
Quartz	30.8%
Spar	33 %
01. 7. 7.	

Chinese	Porcetain	
Clay Substance		31.8%
Quartz		30.8%

German Glaze .11 Potash .67 Lime Aluminum Oxide + 10 Silica .22 Magnesium Oxide

The above glaze is made from

.11 Potash + .11 Aluminum Oxide + .66 Silica = 61.27 lb. Spar .67 Lime = 67 lb. Marble = 18.48 lb. Magnesite

.22 Magnesium Oxide as Magnesite .89 Aluminum Oxide + 1.78 Silica

= 230.5 lb. Zettlitzer Clay 7.56 Silica = 453.6 lb. Hohenbacker Sand

19.4% Natrium Spar Mica 18 %

The firing temperature of the German and Danish porcelain varies between Seger Cone No. 14 and No. 16. household china is usually bisquit fired at a temperature of about 900° C. before glazing and then glazed and given the final firing at about 1400° C. They are then painted or decorated and given short firing at about 600° to set the colors.

The quartz used chiefly in the above mixtures comes from Sweden, the feldspar from Norway, whereas some of the best clays come from Czechoslovakia, although good raw materials are obtainable

in many countries.

The chemical analysis of the above materials is as follows:

Zettlitzer Clay		12.65%
Silicic Acid		46.9 %
Aluminum Oxide		38.56%
Ferric Oxide		.84%
Potash and Sodium	Oxide	1.05%
Giving the followin	g technic	al analysis
Clay Substance	0	98.8 %
Quartz and Spar		1.2 %
Feldspar	Nor-	Czecho-
-	wegian	slovakian
Silicic Acid	62.25%	54.5 %
Aluminum Oxide	19.96%	19.75%
Ferric Oxide	.35%	1.75%
Potash (K ₂ O)	14.32%	11.5 %
Lime (CaO)	.55%	
Magnesia (MgO)	.21%	
Sodium Oxide		
(Na ₂ O)	1.36%	

Magnesia Porcelain

This porcelain usually consists of about 85% powdered talcum and 15% settled gelantinic magnesium silicate, or 80% talcum and 20% China clay. These magnesia porcelains have very good electrical and mechanical properties and extremely small shrinkage during firing allowing the pieces to be made to very close dimensions. They also retain their high electrical resistance up to very high temperatures.

The chemical formulas for some of the glazes used by various European manu-

facturers are as follows:

Another clear and fine German glaze consists of

.3 Potash .8 Aluminum Oxide + 8 Silica

Danish Glaze

.65 Potash Aluminum Oxide + 15 Silica

This glaze is made of

	THIS SIGNED AS INTO OR		
	China Clay	6.75	lb.
(Quartz	48.75	lb.
1	Špar	28	lb.
(Crayon	2.75	lb.
-	Bisquit fired porcelain		

(Powdered) 13.75 lb.

All the above formulas are based on pure European porcelain materials and if materials obtained locally are used a thorough chemical and rational analysis must be made of the raw materials used and the formulas corrected for the varying compositions of the materials.

Low Expansion Borosilicate Glass U. S. Patent 2,012,552

A borosilicate glass having a thermal coefficient of expansion of about .000005 and consisting essentially of silica 72%, magnesia 12%, boric oxide 8%, sodium oxide 6% and potassium oxide 2%.

Ultra Violet Stable Glass British Patent 424,366

Potassium Carbonate	13.77	lb.
Potassium Nitrate	6.71	lb.
Calcium Carbonate	8.93	lb.
Barium Carbonate	3.22	lb.
Magnesium Carbonate	18.53	lb.
Boron Oxide	31.04	1b.
Aluminum Oxide	28.80	1b.
Diammonium Hydrogen		
Phosphate	48.70	1b.

Brown Glass U. S. Patent 2,014,230

A batch for making brown glass comprises in addition to the ordinary glass composition 0.5 to 3.0% of ammonium sulphate and 0.5 to 5.0% of organic matter.

Coloring Glass Austrian Patent 140,547

Colored coatings are produced on sulphide glass not containing free carbon. A typical sulphide glass is made from

-	
Sand	87 lb.
Soda Ash	20 lb.
Source Profit	20 10.

Potassium Carbonate	10 lb.
Lime	11 lb.
Borax	2 lb.
Ferrous Sulphide	3 lb.

Colored Coating Composition

Cuprous Oxid	le	30 lb.
Calcined Copy	per Sulphate	30 lb.
Calcined Clay	7	120 lb.

Luminescent Glass British Patent 415,536

Zinc sulphide and/or cadmium sulphide, etc. are/is either added to the glass or formed in the glass by reduction of the corresponding sulphates with zinc, tin, magnesium powders, carbon, sulphur, etc. or by combination of the oxides or carbonates with sulphur. The presence of 0.01-0.4% of a heavy metal (cadmium, copper, antimony, manganese, etc.) is also necessary. An orange-yellow glass is composed of silicon dioxide 66, aluminum oxide 3, boric anhydride 3, calcium oxide 3, zinc oxide 5, potassium oxide 5.5, sodium oxide 11.5, manganese sulphide 0.63, and zinc sulphide 2.37%.

Cream Colored Opaque Glass U. S. Patent 1,956,176

Fuse together		
Sand	885	lb.
Soda Ash	306	lb.
Feldspar	675	lb.
Cryolite	90	lb.
Calcium Fluoride	50	lb.
Sodium Nitrate	30	lb.
Arsenic Trioxide	4-10	lb.
Ferric Oxide	4-10	lb.
Sodium Uranate	2-7	lb.
Selenium	1/8-5/8	lb.

Vacuum Tube Glass U. S. Patent 1,969,277

	,	,			
Boric Oxide		40	to	60	Ib.
Sodium Oxide		4	to	5	lb.
Calcium Oxide		10	to	11	lb.
Alumina		11	to	13	lb.
Silica		20	to	30	116

Lightly "Frosted" Glass

Gelatin Sodium	Fluoride		$\frac{4.5}{2}$	g.
Water			30	ee.

The gelatin is first dissolved in the water and then the sodium fluoride is added. The solution is then poured over a glass plate and the latter is allowed to dry in a horizontal position. When com-

pletely dry, the plate is immersed in a dilute solution of hydrochloric acid for 30 seconds, and is then again allowed to dry. The remainder of the gelatin may then be removed with the aid of hot water.

Acid- and Waterproof Cement U. S. Patent 1,973,731

Silicate cements are rendered harder and denser by the addition of ½ to 2% of aluminum or calcium hydroxide and sodium silico-fluoride.

Special Cement French Patent 777,055

Portland Cement Clinke	
Slag	50-55 kg.
Silica	8-12 kg.
Slaked Lime	8-12 kg.
Plaster Stone	2-6 kg.
Grind all together.	,

Cellular or Light Weight Concrete U. S. Patent 1,985,905

A slurry is formed by mixing cement with following foam producing compound:

Water	500 lb.
Casein	100 lb.
Slaked Lime	25 lb.
Benzaldehyde	7 lb.
Beta Naphthol	1 lb.
Arsenic Trioxide	1 lb.

Coloring Concrete

For coloring white Portland cement, 5 to 10% of the following materials are generally employed:

Iron Oxides	Red, yellow,
	brown, black
Manganese Dioxide	Brown, black
Chromium Oxide	Green
Ultramarine Blue	Blue
Cobalt Blue	Blue
Carbon Pigments	Black
_	

Certain types of pigments such as those containing Prussian blue, zinc and lead chromates, and cadmium lithopone cannot be used. Chrome green needs to be carefully distinguished from chromic oxide green. Lead oxide pigments are unsuitable and ultramarine is not entirely stable. The fading of colored concretes is due to the formation of a film of calcium carbonate on the surface.

Fire Resistant Concrete Hungarian Patent 109,616

Chamotte Flour (10 mil.gr.)	3 qt.
Cement	1 qt.
Quartz Powder	1 qt.

Waterproofing Mortar and Concrete Austrian Patent 138,387

Olein		10	kg.
Ammonia	(0.910)	2	1.

Mix until uniform and then add slowly with stirring

Aluminum Sulphate (22° Bé.) 2 l. or Zinc Oxide 2 kg

In use, the above mixture is added to 100 times its weight of 20% milk of lime and the latter is used in place of the water to be used with the cement.

Flexible Paving Material U. S. Patent 1,961,678

Approximately 60% of coarse (¼-1½-in.) anthracite bone and rock from a cleaning plant together with fillers, e.g., sand 30 and marble dust 5%, is mixed with 6-12% of a bituminous binder.

Road-Surfacing Material Swedish Patent 80,677

Slabs for road, sidewalk and floor surfacing are made from a mass consisting of 20.4% wood tar, 20.4% coarse sand below 3 mm. size, 40.8% fine sand having a grain size of 0.25-2.0 mm., 4.1% ground unslaked lime, 8.2% cement and 6.1% of fireclay.

Tennis Court and Path Surfacing British Patent 430,001

Twelve pounds rosin are mixed hot with 1 gal. raw linseed oil and 1 oz. powdered alum, 2 gal. of the resulting syrup being mixed with 6 cu. ft. dry sand and 30 oz. chrome green being added. If a quick-setting, tough material is required, 70 oz. of tung oil and 5% (calculated on total oils) of a 4% cobalt linoleate are added.

Asphalt Powder German Patent 613,620

Asphalt (M.P. 45° C.)	3	lb.
Glass or Mica Powder	1	lb.
Warm and mix cool and nor	wder	

Pavement Joint Packing U. S. Patent 2,016,404

Rubber	40	lb.
Asphaltum	7	lb.
Whiting	46	lb.
Sulphur	3	lb.
Ammonium Carbonate	2	lb.
	_	_

Work into a porous mass and cure by heating.

Refractory Compound British Patent 413,398

A mixture of refractory plastic clay with finely ground glass (of any quality) borax and sodium chloride (e.g., 20, 2, 1 and 3 parts by weight respectively) yields refractory products of increased durability and is also suitable for use as a refractory plaster or cement.

Ingot Mold Refractory U. S. Patent 1,984,759

Chrome Ore	8-10 lb.
Basic Slag	25 lb.
Magnesite	10-12 lb.
Calcined Fire Clay	50-30 lb.
Plastic Clay	10-15 lb.
Common Fire Clay	20–28 lb.

Spark Plug Refractory British Patent 422,474

Corundum	96 lb.
Titanium Dioxide	2 lb.
Magnesium Dioxide	2 lb.

Heat; grind; mix with a little acid, mold and fire at 1630° C.

Refractories Resistant to Spalling

Bricks for suspended arches of boiler furnaces can be made of a highly aluminous clay containing silica 54.48, aluminum oxide 43.18, ferric oxide 1.10, calcium oxide 0.86, magnesium oxide 0.18%; ignition loss was 0.32%. No plastic clay was added.

Fused Silica, Improved U. S. Patent 1,984,178

An insulating composition having essentially the properties of fused silica but being characterized by improved workability when plastic and decreased brittleness, consists mainly of silica and contains as constituents about 1/4 to 11/2 per cent of beryllium oxide and about 1/5 to 2% of aluminum oxide.

Inorganic Electric Insulation for Steel U. S. Patent 1,951,039

Steel sheets	are c	oated	with	a mi	xture	,
of						
Lime				15	lb.	
Iron Oxide				28	lb.	
Sodium Sili	cate			70	lb.	
Water				200	lb.	
Bake at 240	° C. a	and an	neal	at 80	0° C	•

Tooth Stump Model for Dental Crowns British Patent 421,872

Aluminum Oxide	50 oz.
Silica	16 oz.
Calcium Sulphate	33 oz.
Gold Chloride Solution (1	l%) 1 oz.

Insulating Decorative Molding British Patent 430,041

Hydrofluorosilic	Acid		15 lb.
Sodium Silicate			8 lb.
Mica Powder			20 lb.
Asbestos			65 lb.
Algolite			15 lb.
Water	to	make	plastic

Sound Absorbing Composition U. S. Patent 1,996,032

Mineral Wool	851/2	1b.
Glue	2	lb.
Cooked Starch	9	lb.
Pyrophyllite	21/2	lb.
Beta Naphthol	1/2	OZ.
Aluminum Sulphate	2	oz.

Treating Peeled Rattan U. S. Patent 1,959,463

The plugs are impregnated with a 1% aqueous solution of glycerol, water is evaporated and the treated plug is sprayed with a solution formed of celluloid 2 lb. and acetone 1 gal. to which powdered aluminum 20 g. and powdered zinc 3 g. have been added, to serve as a sealing and preservative agent.

Minimizing Wood Shrinking and Swelling

Soak wood in water in a vacuum chamber, the air being removed by alternate evacuation and breaking the vacuum. Soak for a week in "Cellosolve" and then distil under vacuum of 60 cm. mercury at 40-45° C. in a number of steps over a period of 3 days. Dry, distil at 100° C. The "Cellosolve" may be sub-

sequently replaced, if desired, by soaking in oil or molten wax for more than a week at temperatures up to 85-90° C.

Wood Antiseptic and Fireproofing British Patent 425,495

Combined fireproofing and preservative properties are claimed for mixtures in aqueous solution of a metallic phosphate, a borate, and a chloride. Impregnation of wood can be undertaken in the usual metal apparatus, since the ingredients are without chemical action on iron. Being resistant to temperatures up to 1000° C., the materials specified not only prevent spread of combustion, but it.

smother flames entirely. These preparations are also said to be suitable for preserving and fireproofing paper, fabrics, etc., by the simple process of soaking. An example of a water-insoluble preparation comprises 5 lb. dibasic sodium phosphate, 3 lb. sodium tetraborate, 1 lb. zinc chloride, 12 lb. 25% aqueous ammonia solution, and 90 pt (maximum) water.

Fireproofing for Wood

Ammonium Phosphate	100	kg.
Boric Acid	10	kg.
Water	1000	1.

Mix and dissolve and immerse wood in it.

Hardness Scale

 Tale Rocksalt Calcite 	4. Fluorite 5. Apatite 6. Feldspar 7. Quartz	8. Topaz 9. Corundum 10. Diamond
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Hardness of Materials

The above numbers give only the order of arrangement as to hardness.

The above	numbers gr	ive only the	oraer	or arrangement	as to naron	ess.
Agate		7.		Hematite		6.
Alabaster		1.7		Hornblende		5.5
Alum		2-2.5		Iridium		6.
Aluminum		2.		Iridosmium		7.
Amber		2-2.5		Iron		4-5.
Andalusite		7.5		Kaolin		1.
Anthracite		2.2		Lead		1.5
Antimony		3.3		Loess (0°)		0.3
Apatite		5.		Magnetite		6.
Aragonite		3.5		Marble		3-4.
Arsenic		3.5		Meerschaum		2-3.
Asbestos		5.		Mica		2.8
Asphalt		1-2.		Opal		4-6.
Augite		6.		Orthoclase		6.
Barite		3.3		Palladium		4.8
Beryl		7.8		Prosphor Bronze		4.
Bell-metal		4.		Platinum		4.3
Bismuth		2.5		Plat-Iridium		6.5
Boric Acid		3.		Pyrite		6.3
Brass		3-4.		Quartz		7.
Calanime		5.		Rock-Salt		2.
Calcite		3.		Ross' Metal		2.5 - 3.0
Copper		2.5-3.		Silver Chloride		1.3
Corundum		9.		Sulphur		1.5 - 2.5
Diamond		10.		Stibnite		2.
Dolomite		3.5-4.		Serpentine		3-4.
Feldspar		6.		Silver		2.5-3.
Flint		7.		Steel		5 - 8.5
Fluorite		4.		Talc		1.
Galena		2.5		Tin		1.5
Garnet		7.		Topaz		8.
Glass		4.5 - 6.5		Tourmaline		7.3
Gold		2.5-3.		Wax (0°)		0.2
Graphite		0.5-1.		Wood's Metal		3.
Gypsum		1.6-2.		\mathbf{Zinc}		2.5

Wood Preservative British Patent 424,941

On impregnating wood with a mixture of a chromate, a salt of a heavy metal—i.e., a metal with a specific gravity greater than 4—and sodium fluoride, a reaction is claimed to take place in contact with the acids and the cellulose in the wood with formation of water-insoluble substances exercising powerful fungicidal action. A preferred mixture comprises 50% potassium or sodium bichromate, 30% zinc chloride, and 20% sodium fluoride, and the impregnation can be effected by standard methods such as a vacuum and pressure process, using a 1% aqueous solution.

Wood Preservative British Patent 425,781

Impregnate with a soluti	ion of
Potassium Dichromate	5.6 lb.
Copper Sulphate	5 lb.
Chromium Basic Acetate	0.53 lb.
Acetic Acid	0.80 lb.
Water	89 lb.

Boric acid and ammonium dihydrogen phosphate may be added for fireproofing.

4	Creosote	Wood	Preservative	Emulsion
	Glue Sulpho	nated Fa	atty Alcohol	0.08 g. 0.02 g.

Creosote	50	g.
Water	50	o.

Allow first two items to swell in water and then mix with crossote and run through colloid mill. Stability is improved by neutralizing any free acidity in crossote with alkali.

Cresylic Wood Impregnati	on Ba	th
Cresylic Acid	100	lb.
Red Oil (Double Pressed)	100	
Caustic Soda Solution 32° I	3é. 20	lb.

Manipulation: Add caustic soda solution to red oil at 50° C., add cresylic acid slowly with constant agitation and cool rapidly.

Arsenic Cement Coating for	Wood Piling
Sand	12 lb.
Cement	3 lb.
Arsenic, White	1 oz.
Mix dry and add water	before use.
Then apply to piling by air	oun.

Oil for Wood Preserva	tion
Carbolineum, Pale (Bleached	l
with Chlorine, Tar Oil)	80 g.
Rosin, Pale	20 g.
Anilin Dye, Oil-Soluble	to suit
Linseed Oil	5-10 g.
Linseed Oil optionally {	1-3 g.

PAPER

Cool to 35° C. and add Ammonia (28°)

2 lb.

Formula No. 1	Ammonia (28°) 2 10.
Argentine or Silver Paper:	Cold Water to make 50 gal.
Argentine Pulp 40% 90 lb. Casein Solution (1/4 lb. per gal.) 21/2 gal. Carnauba Wax Emulsion 1/2 gal. Toluol 1 pt. Carbon Tetrachloride 1 pt.	The emulsion should be allowed to stand for at least 24 hours before use as it seems to improve with age. This emulsion is added to the coating mixture in sufficient amount to give the desired gloss when the paper is flinted.
Nigrosin 9 oz.	No. 4
The casein solution is made as follows:	Canadian Patent 344,222
Casein 62 lb. Borax 7 lb. Trisodium Phosphate 7 lb. Water to make 50 gal. The carnauba wax emulsion is made	Phthalic acid (8.5) and caustic soda (5.5 parts) are dissolved in 1300 parts of water at room temperature. White molding plaster or calcined gypsum (850 parts) is added and the mix is stirred for
with this formula:	1 hour. To this slurry is added 1100
Carnauba Wax 140 lb. Castile Soap 20 lb. Water to make 140 gal. No. 2	parts of casein glue containing 170 parts of dry casein. The product is used directly on the paper-coating machine. The method may be modified for the utilization of a mixture of the deflocculated
A coating mixture which will give a high finish when calendered is made up as follows:	gypsum and coating clay by using soda ash as the electrolyte, and Turkey-red oil may be added to the final product.
Water 65 gal. Soda Ash 3 lb. Ammonia 4 gills Satin White Pulp 440 lb. English Clay 650 lb.	Playing Cards British Patent 405,502
English Clay 650 lb. Stir untill thoroughly mixed and smooth and add the following casein	The cards are composed of a core of textile fabric impregnated with a solu- tion of cellulose derivative and coated on

50 gal.

100 lb.

10 lb.

Trisodium Phosphate	7 lb.
Borax	5 lb.
Ammonia	6 gills
This coating mixture	
high finish when calender	
able for the highest gra	ade lithographic
or process printing.	7 1

solution:

Water

Casein

Soda Ash

Paper Coating

Formula No. 1

No. 3 Wax Emulsion for Flint Paper Yellow Laundry Soap 7 lb. Carnauba Wax 50 lb. Water 121/2 gal.

Boil with live steam till thoroughly emulsified (from 3-4 hours).

Stencil Sheets U. S. Patent 2,004,484

both sides with a layer or layers of cellu-

lose derivative solution containing such

a small amount of plasticizing agents

that the cards are elastic. A suitable

composition consists of cellulose acetate 2.5, acetone 4, denatured alcohol 6, castor oil (plasticizer) 0.35 and dry pigment 0.16 kg. To make the card opaque the composition used for coating one side may contain metallic pigments, e.g.,

Yoshino paper is coated with		
Gelatin	13	oz.
Hard White Soap	42	oz.
Almond Oil	56	OZ.

bronze powder.

Treating Parchment Paper for Wrapping Butter

Parchment for salt butter is immersed for ten minutes in a solution of 2½ lb. salt in 10 gal. water heated to 220° F.

Separating (Non-Sticking) Paper U. S. Patent 2,017,449

A flexible fibrous sheet is coated with Sodium Silicate 140 g. Glycerin 15 g. Carnauba Wax Emulsion 1 g.

Gummed Paper U. S. Patent 1,940,363

A thin film of adhesive composed of 90% of dextrin and 10% of gelatin glue applied to transparent paper enables it to be printed with common quick-drying inks and to adhere to glass.

Waterproofing for Paper

Trihydroxyethylamine
Stearate
Stearic Acid
Water
Steil and mix until smooth; pour into this slowly while stirring vigorously
Paraffin Wax (Heated

to 90-100° C.) 30 lb. Stir until cool.

Use 1 part of above emulsion to 5-10 parts of warm water.

Non-Staining Waterproofing for Paper U. S. Patent 1,968,907

Petrolatum Wax 25–90 lb. Ester Gum 5–75 lb. Paraffin Wax 5–50 lb.

Waterproofing for Paper Australian Patent 5604

Shellac		22	oz.
Alcohol		75	OZ.
Formaldehyde		3	oz.

Waterproofing Paper and Fiber Board Canadian Patent 343,302

The strength and water resistance of fibrous material are increased by beating in a liquor containing 34 to 4 lb. of casein per 100 lb. of pulp lime from 10 to 25% of the weight of the casein, and sodium fluoride from 5 to 12.5% of the weight of the casein. The material treated may be paper, fiber board, as

bestos board or the like. The strength and water resistance may be increased if a relatively small quantity of formaldehyde is added to the treating solution. If the fiber so treated is somewhat too brittle, a softening agent such as glycerol, sulphonated or saponified oil or fat may be added to the treating composition.

Waterproofing Paper and Textiles U. S. Patent 1,981,405

Glue	15 oz	5.
Water	83 oz	Z.
Formaldehyde	1-2 oz	z.

Dissolve glue in water and mix formaldehyde with it vigorously and spray immediately on material to be waterproofed.

Embossed Waterproof Wallpaper U. S. Patent 1,936,355

Stearic Acid	4 lb.
Japan Wax	5 lb.
Triphenyl Phosphate	8 lb.
Dibutyl Phthalate	1 lb.
Heat to 90° C. and add	
Water Shellac (40%)	56 lb.
Triethanolamine	2 lb.
Cool to 70° C. and add	successively
ith vigorous stirring	

Ammonia (28%) 1 qt.
Water 3 gal.
Ammonia (28%) 3 qt.

Water 3 gal. Latex + 4% Sulphur 3 lb. Water to make 28 gal.

Odorless Greaseproof Paper and Textiles British Patent 431,191

This composition comprises a cellulose derivative and rubber or chlorinated rubber dissolved in a solvent free from benzene or its derivatives and containing di-, tri-, or per-chloroethylene and/or methy-lene chloride. The composition may be employed for the production of artificial silk, filaments, threads, films, sheets, and the like, in which case the preferred proportions are chlorinated rubber 30 to 50 parts and cellulose derivative (nitrate or acetate) 800 to 900 parts. A typical solvent for such a mixture comprises trichloroethylene or methylene chloride 180 to 300 parts, and acetone 2000 to 3375 parts. A further application of the composition is in the production of an odorless and grease-proof wrapping paper, and of coated textile and like sheets. A

Alum

suitable composition for this purpose comprises chlorinated rubber 15 to 20 parts, cellulose nitrate or acetate 60 to 80 parts, dissolved in a mixture of trichloroethylene 90 to 120 parts and acetone or methylene chloride 1000 to 1300 parts. To this composition may be added a mixture of diethyl phthalate, castor oil and paraffin oil as plasticizer. A paper base may be coated by passing it through the composition, which is maintained at a temperature of 28-38° C. and the coating dried by passing through a drying chamber. The drying step is preferably followed by a humidifying operation by passing the coated paper through a tower containing humidified air. In place of the cellulose acetate or nitrate there may be used benzyl cellulose. The rubber and the cellulose derivatives may be dissolved together, or may be dissolved separately and the solutions mixed.

Wax Size, Paper Formula No. 1 U. S. Patent 2,009,488

First emulsify a corn oil soap with water to form a paste. Next mix into this paste modified starch in the ratio of preferably approximately about 15 parts of modified starch to 10 parts of corn oil soap. Thereafter, and while the mixture of corn oil soap and modified starch is constantly agitated, incorporate a wax, preferably melted paraffin, although other waxes such as montan, japan, carnauba, etc., may be used alone or in substitution for a portion of the paraffin. The wax may be incorporated in the amount of 75 parts to 15 parts of modified starch and 10 parts of soap.

The mixture thus produced may be incorporated in the beaters in which event add a small percentage of paper manufacturers' alum to aid in the precipitation as the retention of the size is increased in this way. The mixture thus produced may also be used as a surface sizing and so used as mixed with sufficient water to produce the desired fluidity. The amount of water equal to the weight of the wax component is satisfactory.

The corn oil soap prevents foaming in the compounding of the size and the modified starch eliminates to a large degree the softening effect upon the paper heretofore produced through the use of wax emulsion sizes.

The resulting size paper has a high finished hard surface and the sizing is equally applicable to cellulosic and asbestos paper stocks. In connection with asbestos paper, the resulting size renders the paper highly water resistant.

No. 2

Canadian Patent 352,422

Pulp Fiber (Dry Weight)	1,000	lb.
Water	20,000	lb.
Mix in a beater and add		
Calcium Carbonate	300	lb.
Ammonium Resinate		
(Dry Weight)	15	lb.
Water	5 00	lb.

15 lb.

Plant Cover and Fruit Wrapping Paper Canadian Patent 346,222

To each ton of unbleached sulphite pulp is added 160 lb. of thick size, or other suitable size equivalent to 112 lb. of dry size. The stock is beaten for 30 minutes; then 40 lb. of copper sulphate in suitable water solution is added to the stock. Beating is continued for 15-20 minutes. A slight excess of size is maintained with a backwater pH of not less than 6.0. The paper prepared from the stock will contain an excess of the desired 1% per weight of copper resinate: that amount of copper resinate being considered necessary to impart to the paper sufficient resistance to the deterioration and destruction of its fiber when used as a plant cover or fruit wrapper.

Detecting Artificial Watermarks in Paper

Artificial watermarks produced by impression on the nearly dried paper with a rubber stamp are differentiated from the genuine by sprinkling the area with a mixture of 100 g. of dry icing sugar and 0.5 g. of concentrated Rhodamine-6G, placing the paper in a dish of water, and examining in filtered ultra-violet light. The design of genuine watermarks is marked for a few seconds by a bright golden fluorescence, which is absent in the case of artificial watermarks.

Discharge Effects on Tissue Paper

Discharge effects on tissue paper are produced in a very simple manner by passing the tissue paper through the solution of an easily dischargeable dyestuff in the dyeing machine, and spraying or printing on a solution of 1 lb. Hydralite C extra per 1 gal. water to which has been added a solution of 3½ oz. acetate of zinc per 1 gal. water or 1 pint acetate

of alumina of 18° Tw.; the paper is then dried quickly.

Increasing Strength of Paper U. S. Patent 1,997,487

An absorbent	paper	is	treat	ed	with
Glue				6	oz.
Formaldehyde				3	fl. oz.
Water	t	o	make	1	gal.

Transfer Printing Paper U. S. Patent 1,965,257

Rubber Latex (60%)	40 lb.
Casein	10 lb.
Zinc Stearate	5 lb.
Paraffin Emulsion	50-100 lb.
Formaldehyde (40%)	5 lb.
Triethanolamine	2 lb.
Water	2 lb.

The colored design is printed on this paper by using a dye ink having a composition similar to the following:

Acid Dye	100 lb.
Acetone	300 lb.
Divinyl Resin	150 lb.
Methyl Alcohol	100 lb.
Dibutyl Phthalate	10 lb.
Castor Oil	10 lb.

To assist the transfer of the colored pattern to the silk fabric it is advantageous to have present at the time of pressing a volatile solvent which is capable of dissolving the dye but not the coating composition. For assisting the transference of acid dyes to silk fabric it is found that a satisfactory solvent consists of:

Alcohol (95%) Acetic Acid (36%) Water	80 gal. 10 gal.
water	10 gal.

It is claimed that owing to the resiliency of the rubber coating composition and the special manner of applying the transfer paper to the silk fabric, it is possible to obtain very clear and well-graded impressions on crepe materials.

PHOTOGRAPHY

Fixing Baths	A fresh bath should be prepared fre-
Acid Fixing Bath	quently, as the gelatin-coated backs of
Metric Avoirdupois	the films are likely to become stained in
Water 4 l. 128 oz.	an old or discolored fixing solution. The
Hypo 1160 g. 38 oz.	Tollowing Replenisher for two-liter solu-
Potassium Meta-	tion of above fixing bath is recommended
bisulphite 100 g. $3\frac{1}{2}$ oz.	in cases where the acidity needs to be
The metabisulphite should be added	renewed:
only when the hypo solution is cool, not	Metric Avoirdupois
when it is hot.	Water 80 cc. 3 oz.
Chrome Alum Fixing Bath	Sodium Sulphite
Solution 1	(Anhydrous) 15 g. ½ oz. Acetic Acid
Metric Avoirdupois	(28% Pure) 48 cc. 1½ oz.
	Potassium Alum 15 g. ½ oz.
Water 2½ l. 80 oz. Hypo 960 g. 2 lb.	Special Fixing Bath for Printon and
Sodium Sulphite	Reprolith Films
	Accuracy in registration for multi-
(Anhydrous) 65 g. 21/4 oz. Water to make 3 l. 96 oz.	color work being of prime importance
Solution 2	101 use in such cases a fixing hath with.
Metric Avoirdupois	out hardener, as follows is recommended:
Water (About	Metric Avoirdupois
Water (About 150° F.) 1 l. 32 oz.	Water 1 l. 32 oz.
Potassium Chrome	
Alum 60 g. 2 oz.	Potassium Meta-
Sulphuric Acid C.P. 9 cc. 1/4 oz.	bisulphite 75 g. 2½ oz.
Add solution 2 slowly with constant	In case this bath should lose its acidity
stirring to solution 1.	by frequent use, giving the film a vellow-
Acid Hardening Fixing Bath	ish stain, add more potassium metabisul-
Solution 1	phite to restore the acidity of the solu-
Metric Avoirdupois	
Water 4 l. 128 oz.	Acid Translation Et :
Water 4 l. 128 oz. Hypo 960 g. 2 lb.	Acid Hardening Fixing Bath
Solution 2	U. S. Patent 1,981,391
The state of the s	Formula No. 1
Metric Avoirdupois Water (About	Sodium Thiosulphate 300 g.
125°F.) 300 cc. 10 oz.	Sodium Sulphite (Desic-
Sodium Sulphite	cated) 15 g. Propionic Acid 20 g
(Anhydrous) 60 g. 2 oz.	
Acetic Acid	
(28%) 180 cc. 6 oz.	Water 5 g.
Potassium Alum 60 g. 2 oz.	No. 2
To make 28% acetic acid from glacial	Sodium Thiosulphate 300 g.
acid, dilute 3 parts glacial with eight	Sodium Sulphite
parts of water.	(Desiccated) 15 g.
Dissolve chemicals thoroughly in order given. Cool solution 2 after mixing and	Acetic Acid 15 cc.
add it slowly with constant stirring to	Potassium Alum 15 g.
solution 1.	Western Borate 10 g.
950	to 1 h.

	PHOTO
No. 3	
Sodium Thiosulphate	300 g.
Sodium Sulphite	500 8.
(Desiccated)	15 g.
Boron Triacetate	15 g.
Potassium Alum	15 g.
Water	to 1 1.
No. 4	0.00
Sodium Thiosulphate Sodium Sulphite	300 g.
Acetic Acid	15 g. 15 cc.
Acetic Acid Citric Acid	15 cc. 1 g.
Potassium Alum	15 g.
Boric Acid	5 g.
Water	to 1 I.
No. 5	
Sodium Thiosulphate	300 g.
Sodium Sulphite	5 g.
Acetic Acid Sodium Acetate	20 cc. 20 g.
Potassium Alum	00.0
Borax	20 g. 20 g.
Water	to 1 l.
No. 6	
Sodium Thiosulphate	300 g.
Sodium Sulphite (An-	
hydrous)	15 g.
Sodium Acetate (An-	20 g.
hydrous) Boric Acid	
Sulphuric Acid (Con-	5 g.
centrated)	5 cc.
Alum	15 g.
Water	to 1 l.
, 2000	
Elon-Hydroquinone De	veloper
Stock Solution	
Avoirdupe	
Elon 45 gr	r. 3.1 g.
Sodium Sulphite	
(Desiccated) 1½ or	10 0
Hydroquinone 175 gr Sodium Carbonate	r. 12 g.
(Desiccated) 2 ¹ / ₄ oz	z. 67.5 g.
Potassium Bro-	
mide 27 gr	r. 1.9 g. z. 1 l.
Water to make 32 oz	
Dilute 1 part to 2 parts w	ater for use.
Di anvilandiamina Da	lanau
p-Phenylenediamine De	veloper
p-Phenylene-	. 10 ~
diamine 145 gr Sodium Sulphite	. 10 g.
(Desiccated) 1 oz	. 50 g.
290 gr	0
Water to make 32 oz	
TO 1 1 1: C1 :	D
p-Phenylenediamine-Glycin	Developer
p-Phenylene-	. 10 g.
diamine 145 gr Glycin 175 gr	40
Cracin Tio St	. 12 g.

GRAPHY	259
Sodium Sulphite (Desiccated) 3 oz. 90 Water to make 32 oz. 1	g. l.
Fine Grain Developer	
Formula No. 1	
Avoirdupois Me	tric
Elon 29 gr. 2 Sodium Sulphite	g.
(Desiccated) 3 oz. 100 145 gr.	g.
Hydroquinone 73 gr. 5 Borax (Crystals) 29 gr. 2 Water to make 32 oz. 1	g. g. l.
No. 2	
Sodium Sulphite 60 p-Phenylenediamine 10 Acetone 10 Sodium Metasilicate 3 Metol 2 Glycin 2 Water to make 2	g. cc. g. g. g.
This is developed for 15 minute 65° to 70° F.	s a.

Pyrocatechol Develor Sulphite	per withou	t
a. Pyrocatechol Water Lactic Acid	4 g. 100 cc 10 dr	
For contrasty negatives	s use	-
a (Above)		ec.
Water	100	cc.
Sodium Carbonate So-		
lution (3-4%)	5	cc.

Developer for	Film	and	Pape	r
Adurol				gr.
Sodium Sulphite Sodium Carbonate				gr.
Water	е		8	gr.

Add not more than ½ grain potassium bromide to each ounce of finished de-veloper. With developer at 70° F., films will develop in 4 minutes. Tuma Gas paper should be exposed so that the image will appear in 45 seconds. The print will be fully developed in 2 minutes.

The times for other papers are:

Velour black-image appears in 1 minute; developed in 21/2 minutes.

Bromide papers-image appears in 11/4 minutes; developed in 3 minutes.

Warmer tones can be obtained by diluting the developer and giving longer exposure.

This developer will not affect persons subject to aniline poisoning. It oxidizes quite rapidly and should be kept in a tall, narrow vessel between prints in order to reduce the amount in contact with the air to a minimum.

Gold toner:

Stock Solution

1. Gold Chloride 15 gr. Water 2 oz.

2. 5% Solution Thiourea

(1 oz. to 20 oz. water) For use take 4 drams of gold solution, drams thiourea solution, 5 or 6 drops

3 drams thiourea solution, 5 or 6 drops sulphuric acid and one quart of water. Proceed as follows:

Dilute the required amounts of both stock solutions with one pint of water. Pour gold solution into thiourea solution slowly with stirring. Add the acid to the combined solutions.

Compensating Developer with Pyrogallol

Formula No. 1		
Water	100 c	c.
Pyrogallol	0.3 g	; .
Potassium Metabisulphite		
(10%)	3 c	c.
Caustic Soda (10%)	2 c	с.
No. 2		
Water	100 c	
Pyrogallol	0.3 g	
Potassium Metabisulphite		
(10%)	12 c	c.
Courtie Sode (100%)	5 0	0

Caustic Soda (10%) 5 cc. Formula No. 1 at 18° C. (5 to 6 minutes) gives a yellow-brown negative.

Formula No. 2 at 18° C. (10 to 12 minutes) gives a neutral gray negative and developer can be used a second time.

Modified Hub No. 1 Formula for Glycerin Developer

		cc.	or	(1	qt.)	
Sodium Sulphite	75	g.		$(2\frac{1}{2})$		
	25	g.		(375)	oz.)	
Trisodium Phos-						

phate (Mono-hydrate) 125 g. $(4\frac{1}{6} \text{ oz.})$

Potassium Bromide 3 g. (45 gr.)

This stock solution keeps well, even in partially filled bottles. For use with chloride and chloro-bromide papers it is diluted with 3 parts of water, and with 4 parts of water for bromide papers. With bromide papers it has been successfully used at temperatures up to 90° F. Because of its high alkalinity, prints developed in this formula should be left in the acid-stop bath for at least 15 or 20 seconds before being placed in

the fixer, and the acid-stop bath should be frequently renewed.

Farmer's Reducer

In case of overexposure or overdevelopment, this well-known reducer can be used effectively for clearing. It is easily compounded by making first a 1:4 solution of plain hypo-for example, 8 oz. of hypo dissolved in 32 oz. of waterand adding to this just enough potassium ferricyanide to turn the solution to a lemon-yellow color. Most workers prepare the ferricyanide as a 10% solution in advance, for use as needed; others shake a little of the powder directly into The lemonthe plain hypo solution. yellow color disappears with use of the reducer, but may be restored by adding more ferricyanide. The stronger the color, the stronger the reducing action, and vice versa. If the reducer is used too strong its action is not so easy to control.

The film may be immersed in the reducer solution, after being soaked in water to assure even action, or, in cases where only local reduction is desired, the reducer may be applied to the moist film with a tuft of cotton, with rinsing during

inspection and afterwards.

Reversing Reversible Film

(1) First Developer

	Metr	ic .	Avoird	lupoi
Water	1000	cc.	. 32	oz.
Metol		g.		gr.
Sodium Sulphite		_		•
(Anhydrous)	30	g.		OZ.
Hydroquinone	12		180	gr.
Potassium Bromide	8	g.		
Sodium Hydroxide	18	g.	1/2	OZ.
Potassium Sulpho-	_			
cyanate		g.		
Develop 4 to 6 mi	nutes	at	65° I	r., de
pending on exposure.				

(2) Wash 5 minutes in running water.

(3) Reversing Bath

Water	$1000 \ cc.$
Potassium Bichromate	5 g.
Sulphuric Acid (Con-	_
centrated)	5 cc.

Normal bleaching time 3 to 6 minutes. Keep in bleaching bath until negative image is completely dissolved.

(4) Wash 5 minutes in running water

(5) Clearing Bath

Water	1000 cc.
Sodium Sulphite (Dry)	50 g.
Clear for 5 minutes.	

(6) Wash 5 minutes in running water.

(7) Expose to Mazda light or diffused daylight.

(8) Second Developer.

Water	1000	cc.
Metol	5	g.
Hydroquinone	6	g.
Sodium Sulphite (Dry)	40	g.
Potassium Carbonate	40	g.
Potassium Bromide	6	g.
Develop 5 minutes at 65°	F.	

(9) Short rinse in running water.

(10) Fixing Bath

, 0	
Water	1000 cc
Нуро	300 g.
Potassium Metabisulph	ite $50 \mathrm{g}$.
Fix for 2 minutes.	•

(11) Wash for 30 minutes in running water.

(12) Glycerine Bath

1000 cc. Water Glycerin (C.P.) 20 cc.

Leave in glycerin bath for 5 minutes.

(13) Remove water with a soft chamois and dry in a current of warm dry

Note: Operations 7 to 13 take place in white light.

Superpan Reversible film can be desensitized before development by immersion in a 1/5000 solution of Pinacryptol Green desensitizer.

Formula "D16" for Chemically Reversing 16 mm. Film

versing 10 mm.		TITIL		
Water (Distilled)			10	gal.
Elon		I	180	gr.
Sodium Sulphite	3	lb.	5	oz.
Hydroquinone			8	OZ.
Sodium Carbonate		lb.		
Potassium Bromide	1	oz.		
Citric Acid			100	gr.
Potassium Metabisulph	ite	3	2	oz.
Develop 7-15 minutes	$^{\mathrm{at}}$	65°	F.	

Intensifying Formulas

On some occasions and for certain types of work it may be found desirable to intensify film negatives. In such instances the following formulas will give best results, being desirable for their freedom from stain as well as their effective intensifying action.

Mercury Intensifier:

	Metric	Avoirdupoi
Water	1 l.	32 oz.
Mercuric Chloride	10 g.	150 gr.
Potassium Bromide	5 g.	75 gr.

Chromium Intensifier:

This formula gives slightly more vigorous intensification than the Mercury Intensifier above. Prolonged intensification with it, however, leaves the film with a slight yellow color.

Metric Avoirdupois Water 1 1. 32oz. Potassium Bi-

chromate 9 g. 135 Hydrochloric Acid 6 cc. 1.6 dr.

Immerse negatives in this solution until bleached, wash for 5 minutes in running water, and redevelop in a Metol Hydroquinone developer. The negatives should then be given a 15-minute wash

before drying.

Some intensifying solutions have been known to cause a slight blue coloration of the base of the film. While this is not harmful and does not prolong the printing time unduly, if preferred, such coloration may be easily removed as outlined in the formula for Washing and Drying.

Monckhoven's Intensifier:

Solution A		Avoir-
	Metric	dupois
Water	1 l.	32 oz.
Potassium Bromide	23 g.	3/4 oz.
Mercuric Chloride	23 g.	3/4 oz.
Solution B	_	Avoir-
	Metric	dupois
Water	1 l.	32 oz.
Potassium Cyanide	23 g.	3/4 oz.
Silver Nitrate	23 g.	3/4 oz.

The silver and the cyanide are dissolved in separate lots of water, and the former added to the latter until a permanent precipitate is produced. The mixture is allowed to stand 15 minutes, and after filtering, forms Solution B.

Place the negative in A until bleached through; then rinse and place in Solution B. If intensification is carried too far, the negative may be reduced with a weak solution of hypo.

Because of the deadly poisonous character of this intensifier, it should be used with care and bottles containing it should be suitably marked.

Agfa Mercuric Iodide Intensifier:

Metric Avoirdupois 200 to 20 to 300 cc. 30 Water oz. Mercuric Chloride 100 cc. 10 (2%)Potassium Iodide 25 cc. 2.5 oz. (10%)Hypo (10%) 40 cc. 4 oz. Part of the mercury solution is added to the water and then part of the iodide solution, continuing until all the mercury and iodide is added to the water.

When solution is clear, add the hypo.

Use full strength.

Mercury Intensifier

This is a satisfactory two-solution intensifier for increasing the printing density of thin, flat negatives. This intensifier has the advantage of not staining negatives as readily as other intensifiers when traces of fixing solution have not been completely removed in washing:

Water 1 l. 32 oz.

Mercuric Chloride 40 g. 1½ oz.

Solution B:

Metric Avoirdupois
Water 1 l. 32 oz.
Potassium Iodide 100 g. 31% oz.

Add B to A until the solution clears. Negatives are immersed until changed to a brown color, then washed and redeveloped in a metol-hydroquinone developer such as Agfa No. 64. The intensified negatives need not be fixed, but should be given a 15-minute wash before hanging them up to dry.

Intensifier, Photographic

Mercuric Chloride Solution (20%) 1 fl. oz.

Potassium Iodide Solution (5%) 1 fl. oz.

Sodium Acetate Solution (7%) 1 fl. oz.

Intensifier for Very Weak Negatives
Water 400 cc.
Mercuric Chloride 2 g.
Potassium Iodide 6 g.

Each of the dry ingredients is dissolved in one-half of the water and the two solutions are then mixed. A red precipitate will form at first but will again dissolve, a clear solution resulting.

While the negative attains considerable and rapid intensification, it becomes badly colored and will not last very long. To avoid this, the negative is placed in a solution of sodium sulphite for a period of ½ to 2 hours. It is then washed thoroughly in water.

If the intensification should be too great it may be reduced in a solution of

sodium cyanide.

Toning Formulas

Sepia Tones by Redevelopment:

Sepia tones may be obtained in any print by subsequent treatment after the print is ordinarily finished. The print should be thoroughly washed before treatment to produce a sepia tone. It is then immersed in the bleaching bath (Solution No. 1) for about 1 minute, or until the middle tones of the print are just perceptible. It is next rinsed thoroughly in cold water and transferred to the redeveloper. When original detail has returned and the print is of desired strength (this will take about half a minute), remove print, rinse thoroughly, and harden by immersion for 5 minutes in the Hardening Solution specified for use in connection with the Fixing Bath Hardening Solution only-no Finally, remove the print and (the Hypo). wash for 30 minutes in running water.

No. 1—Stock Solution

(Bleacher)

The No. 1 Stock Solution, which is the bleacher, may be made up for either normal sepia tones, warm sepia tones, or cold sepia tones, as follows:

For Normal Sepia Tones:

Metric Avoirdupois Potassium Ferricyanide (10% 500 cc. Solution) 16 OZ. Potassium Bromide (10% Solution) 100 cc. 31/2 oz. Water 400 cc. 14 OZ.

For Warm Sepia Tones:

Metric Avoirdupois
Potassium Ferricyanide (10%
Solution) 600 cc. 19½ oz.
Potassium Bromide (10%
Solution) 40 cc. 1½ oz.
Water 360 cc. 12 oz.

For approximately a 10% solution, take 100 grains to 2 fluid ounces of water or 10 grams to 100 cc. of water.

For Cold Sepia Tones:

	Met	ric	Avoird	innoi	a
Potassium Ferri-			22,0110	upor	13
cyanide (10%					
Solution	300	cc.	10	oz.	
Potassium Bro-					
mide (10%					
mide (10% Solution)	500	cc.	16	oz.	
Ammonia (.910)	10	cc.	1/2	oz.	
Water	190	cc.	$6\frac{1}{2}$	oz.	

No. 2—Stock Solution (Re-Developer)

Metric Avoirdupois
Water 500 cc. 16 oz.
Sodium Sulphide 42.5 g. 1½ oz.
Bleaching Bath for Use.

Water Solution (Bleacher) Svoir-dupois (Bvoir-dupois 16 oz. 16 oz. 16 oz. 16 oz.

(Bleacher) 500 cc. 16 c Re-Developing Bath for Use.

> Avoir-Metric dupois

Water 1 l. 32 oz. No. 2 Stock Solution (Re-Developer) 118 cc. 4 oz.

Important: Be sure to use sodium sulphide, not sodium sulphite, in compounding the re-developer. Also, use clean trays, free from exposed iron spots, especially with Bleaching Bath. Otherwise blue spots may form on prints.

1. Blue Toner (Iron Bath)

First dissolve:

Potassium Ferricyanide 375 g.
Potassium Bichromate ½ g.
in:

Water 40 1. and pour this solution into a second one consisting of:

Iron Ammonia Alum425 g.Oxalic Acid500 g.Water40 l.

The two solutions must be separately filtered and then mixed at ordinary temperature and with vigorous stirring. They then form a clear yellowish solution without any sign of turbidity, provided the chemicals have been mixed in the correct quantities and with due regard to cleanliness. The time of toning varies according to the tone required.

By treating the toned films in a subsequent weak fixing bath, tones of remarkable clearness are obtained. But it must be expressly noted that, in the case of blue-toned films, the fixing bath must not be used until after a most thorough washing, otherwise a reducing action takes place and detail in the picture is eaten out. The films must be well washed after the second fixing.

2. Uranium Toner (Yellow-Brown)
Dissolve:

Potassium Ferricyanide 500 g. in:
Water 10 l.

and add:

Potassium Bichromate
(1% Solution)

Then add the whole to:

Uranium Nitrate
Oxalic Acid

50 cc.

50 g.

500 g.

Water

As in the making up of the iron bath, the two solutions must be separately filtered and mixed at ordinary temperature while stirring well. The result should be a yellowish solution free from deposit. The bath requires to be revivified from time to time during long use by addition of oxalic acid. As much as 1000 g. oxalic acid may be added in all, and so 500 g. of the acid is dissolved in water, and the solution added in small doses from time to time. By this means, staining of the whites, which otherwise takes place after a time, is readily avoided.

3. Copper Toning (Reddish-Brown)
Copper Sulphate 500 g.
Sodium Citrate 2500 g.
Or
Potassium Citrate 2250 g.
Water 30 l.

To the above add:
Potassium Ferricyanide 400 g.
Water 20 l.
Potassium Bichromate (1% Solution) 50 cc.

In making up this bath also, the separate solutions must be filtered, and carefully and well mixed at the ordinary temperature.

The following observations apply to the use of Baths Nos. 1 to 3.

As is well known, a solution of potassium ferricyanide in water, in conjunction with hypo, acts as a reducer, viz.: the Farmer reducer. Thus a film which contains only traces of hypo, on being introduced into the toning bath, undergoes a reducing process along with the toning which is aimed at. There are also two conditions which should be invariably observed if it is desired to carry out the toning processes successfully and to keep the toning baths in good conditions:

1. For all toning processes—and this applies also to tinting—frames should be kept for these operations only; frames which have been employed for the development or fixation of prints should on no account be used.

2. Positive film which is to be toned must be especially well washed. In order to ensure that this is the case and to be certain that the film is in the necessary state of uniformity, it is advisable to wash the film for a further few minutes immediately before toning.

Wet Collodion Continuous-Tone Negative
Plain Collodion 10 g.
To the above add 1 g. of following:
Alcohol 1 l.
Cadmium Iodide 80 g.
Ammonium Iodide 40 g.
Cadmium Bromide 10 g.
Calcium Chloride (6H₂O) 10 g.

Re-development increases the opacity if done before fixing and increases the contrast if done after fixing.

Prevention of Haze in Prints German Patent 594,712

The formation of haze is prevented and a blue-black tone imparted to the prints, by adding triazole or tetrazole solution to the emulsion layer or to the developer. Thus, 0.5 to 5 cc. of a 1:100 benzotriazole solution is added to a usual metol-hydroquinone developer.

Control of Photographic Contrasts M.

Potash Metabisulphite	160 gr.
Metol	160 gr.
Soda Sulphite	3/4 oz.
Potash Bromide	25 gr.
Water	to 10 oz.
Q.	
Potash Metabisulphite	160 gr.
Hydroquinone	160 gr.
Soda Sulphite	3/4 oz.
Potash Bromide	40 gr.
Water	to 10 oz.
A.	
Soda Carbonate	6 oz.
Water	to 20 oz.

These are concentrated solutions that will keep indefinitely if properly compounded and are diluted for use. In the M and Q solutions the potash metabisulphite should be added to about three-fourths of the water first and partially dissolved, it is not necessary that it should be fully dissolved at this stage, just a good shake up to drive off the oxygen from the water, then the metol or hydroquinone added and fully dissolved before the soda sulphite is added.

For use the M and Q solutions are used either separately or in any proportion desired and an equal volume of the A solution added and then diluted with 3 times the volume of water. For example, for a normal developer take 1 part of M, 4 parts of Q and 5 parts of A diluted with 15 parts of water. The quantity of water can be varied to suit the particular brand of plate in use, some plates will stand twice this quantity of water. It is a matter of experience.

For positives from very flat negatives the Q solution plus A may be used alone, or a small quantity of M such as 1 of M to 10 or 12 of Q. From very hard negatives the M plus A alone can be used or with a small proportion of Q and, of course, the necessary dilution in each

With a high proportion of M to Q the image will appear quickly, but will require time to gain sufficient density. With a high proportion of Q to M the image will appear slowly, but gain density more rapidly in proportion so that the total developing time does not vary so much as would appear at first sight.

To those who have to handle this class of work, either for color half-tone or for photogravure, this system of working is recommended and when once mastered it becomes a very adaptable servant.

Re-Etching Half-Tones with Enamel Off

As in all etching, cleanliness and freedom from grease in the plate to be treated is the first consideration, but any enamel still remaining on the dots is to be left. (This applies to the places to be rolled as well as those where the enamel is good.)

A viscid solution of gum and process white is next prepared:

Gum Arabic 5 oz.
Water 5 oz.

and when required, to every three parts of this solution, mix one part process white.

The plate after being rinsed with water to replace the air between the dots is allowed to drain (not dry) and the gum solution painted over the whole sur-The edge of a wooden rule is next wiped or scraped over the surface in such a way that only the thinnest layer of gum is left on top of the dots leaving the thick gum remaining between them. A word of warning-should the gum become somewhat thin owing to its application to a wet plate the process must be repeated. Also do not put the gum on a dry plate as it would then be impossible for it to replace the air between the dots. After applying the gum it is dried, using as little heat as possible.

A piece of charcoal having on one of its sides a perfectly flat area of about 1 inch, is now required for rubbing the gum off the surface of the plate, and must be used dry. This flat side is put in contact with the gummed surface and with an even and gentle pressure the gum is rubbed away from the whole surface, or if only to be treated locally, from those parts which are to receive the new ink top. It will be found that very little rubbing is required to remove the gum in the high-lights, while this increases somewhat with the strength of the tone. Rubbing is continued until the metal appears bright and clean, removing any enamel that remains on the areas to be rolled at the same time. If this is carried out properly any increase in tone values owing to the rubbing of the charcoal is negligible, and cannot be seen on a graded strip although etched down beside enamel receiving identical treatment. The gum is now remaining at the sides and between the dots untouched, and the powdered charcoal must be lightly dusted off the surface with cotton wool.

It will be noticed that the white gum between the dots is discolored by the charcoal but this does not matter as in other respects it is quite unaffected. At this stage the roller and ink must come under consideration and these contrary to the usual rule are quite easy to prepare and use. The roller used is a good quality composition roller, and the ink is stone to stone re-transfer ink, both ink and roller are the same as used by line metal printers. Thin the ink with a little pure turpentine in the center of the slab and then evenly distribute the ink over roller and slab. The amount of ink when ready for rolling should be such that it is still possible to see the color of the slab through the ink. The condition of the ink should be just tacky. In rolling up the plates no extra pressure is required, the weight of the roller itself usually being found sufficient. When the whole surface of the plate has received an even layer of ink it is dusted over with fine bitumen powder. This dusting must be done lightly and thoroughly with the aid of cotton wool.

The plate should now be soaked in water for about 2 minutes to soften the gum, but soaking only will not bring it away from between the dots, as a certain amount of force is necessary in the form of a spray of water. The spraying can be done by turning the tap on full and putting the thumb in a position so as to make the water into a narrow beam of as much force as possible, and this is directed all over the surface of the plate, dwelling particularly on those parts (if any) where the gum appears somewhat reluctant to leave, such as a strong cross-Should any difficulty be exline tint. perienced in cleaning away ink-covered gum from between the dots, the fault can usually be traced to the gum solution being too thin, or to its imperfect application, but in any case do not attempt other means of removing the gum, such as rubbing with cotton wool, as this will certainly weaken the new top.

After spraying the plate is drained and dried off over the gas with gentle heat, making sure that all moisture is removed before burning-in hard. The temperature reached during the fusing or burning-in of the bitumen and ink should be almost sufficient to burn-in enamel. The required temperature can be judged quite easily in copper by the discoloration of the metal: it turning from an orange to a bluish color when approximately the temperature is reached. In zinc there is no discoloration of the metal, but one way to assist the judgment is to paint the back of the plate with shellac and when during the burning-in this turns a dark brown shade, the ink is burnt in.

Burning-in operations completed, the plate, either copper or zinc, is ready for etching as soon as it becomes cold, and it can be chalked with magnesia, staged and treated as though the dots had the original enamel top. One precaution is necessary and that is to take care they are not immersed for any length of time in the acetic and salt bath other than that required to remove the magnesia, as this has a weakening effect on the ink. It is better to dispense with acetic and use a weak solution of nitric acid such as 1 part acid to 20 parts water.

When etching is completed it is sometimes found difficult to remove the ink top even though turpentine and a brush is used, in which case a light rubbing with charcoal will be found the most

satisfactory.

Photolithographic Deep-Etched Plates

A fine-grained zinc plate is washed with 5% acetic acid and water, then coated with 1000 cc. of water, 133 cc. of photo-engraver's glue, 100 cc. of 20% ammonium bichromate solution, 20 cc. of ammonium hydroxide at 22° Bé. At 30% relative humidity, the exposure is twice as long as at 60%. The sensitized plate keeps 6 hours at 45 to 50% humidity or 24 hours at 40%. After development in

cold water, the plate is treated for 10 to 15 seconds in hydrochloric acid diluted with 200 parts of water, washed, and dried. Before drying, the image may be dyed in a 2% solution of direct black 2N extra concentrated, or oxydiazol black NJEE. Etching the plate in denatured absolute alcohol to which are added 50 cc. of concentrated hydrochloric acid per liter, for 2 minutes, produces a depth of about 0.0075 mm. The plate is rinsed with alcohol, dried, washed out with asphaltum and liquid reversing ink, and talcked. It is then swabbed with water and in 1000 cc. of water, 400 cc. of 10% barium chloride solution, and 50 cc. of 10% sodium hydroxide solution. Removal of the glue image takes from 5 to 10 minutes. This batch is patented in the United States: 60 cc. of 12 to 14° Bé. Gum arabic solution may be added. After washing with water the plate is bathed for 10 to 15 seconds in very dilute hydrochloric acid, then rinsed in hot water. The plate is next gum-etched and sent to the press.

Photoengraving Enamel U. S. Patent 2,000,453

• Glue 20 oz., ammonia solution 2 oz., chromic acid 1.5 oz., and alcohol about 64 oz. are used together.

Planographic and Offset Plates British Patent 421,217

Aluminum plates are made anodes in 0.3-5% nitric for 10 to 30 minutes at a current density of 1 to 2 amperes per square decimeter; zinc plates are made the anode in a saturated potassium carbonate solution for 10 to 30 minutes at a current density of 2 to 3 amperes per square decimeter.

Photographic Masking Paste

Glycerin			1 gal.
Whiting			 3 lb.
Neutral	Soft	Soan	1 lb.

Masking paste must be so formulated as to have sufficient solids or bodying agents that it will not flow down or cause breaks in the film; also it must be capable of being brushed on to form a clean sharp edge. The proportion of glycerin must be sufficient to keep the film from drying up under exposure for at least 48 hours.

Photograph Paste

Gelatin (Photo) 4 oz. Water 16 oz.

Soak, dissolve on a water-bath, and add when somewhat cooled:

Glycerin 1 oz. Wood Alcohol 5 oz. Mix.

Mounting Translite Prints on Glass

Dissolve 1 oz. of gelatin in 6 oz. of boiled water. After the gelatin has been thoroughly dissolved, add 1 oz. chloral hydrate. Apply the solution to the glass with a brush, coating the glass evenly. Then apply Translite print, wet, face side to the glass. Squeegee with a print-roller until all the surplus gelatin has been removed and air-bubbles are all out. Then allow to dry. This formula will withstand heat more than any other starch or glue formulæ.

Photographic Dry-Mounting Tissue U. S. Patent 2,017,144

A paper mounting tissue is coated on both sides with a composition containing low-viscosity nitrocellulose 100, tritolyl phosphate 110-150 and a resin such as shellac 10-200 parts.

Blue for Drawings

Saturate 10 g. of oxalic acid in a little water with ferric hydroxide, filter off excess of ferric hydroxide, add concentrated solutions of 27 g. sodium oxalate and 11.6 g. sodium ferrocyanide, apply the mixture to paper with a brush and dry in a dark room. Develop the prints with dilute hydrochloric acid or sulphuric acid

Waterproof Coating for Wooden Photographic Trays

Formula No. 1

Methyl Alcohol	500 cc.
Orange Shellac	100 g.
Rosin	25 g.
Venice Turpentine	25 g.

The ingredients are heated on a water-bath until completely dissolved.

No. 2

One part of gutta percha and one part of paraffin are melted together. When cool, this mixture is dissolved in sufficient benzine to make a mixture of paint-like consistency.

Cleaning Porcelain Photographic Trays
Water 100 cc.
Potassium Cyanide 10 g.
Iodine 3 g.

This is a very satisfactory solution for removing stubborn stains.

Flashlight Powder Formula No. 1

Potassium Chlorate 20 g. Powdered Magnesium 10 g.

The potassium chlorate must first be finely pulverized (to avoid spattering on ignition). It is then carefully mixed with the magnesium. It is preferable to mix this in small quantities on a glass plate, as this mixture is very explosive and a pestle and mortar may prove extremely dangerous.

No. 2

Powdered Magnesium 10 g. Potassium Dichromate 10 g.

This powder is designed to burn from 1/4 to 3/4 second.

No. 3

Powdered Magnesium 1 g. Ammonium Nitrate 0.8 g.

The above should be mixed just before using, the ammonium nitrate being kept in an absolutely dry state. This is a very brilliant and ashless powder and the quantity designated is sufficient for good illumination of a room 15 ft. sq.

Magnesium Flashlight Powder German Patent 592,898

Potassium permanganate, potassium nitrate and sulphur are among the ingredients of a new type of magnesium flashlight powder composition which can be ignited without detonation in cartridges through the medium of a percussion cap. 700 to 900 parts of magnesium are admixed with sulphur (10 to 18), potassium permanganate (100 to 140), potassium nitrate (70 to 85), magnesia (100 to 160) and wood charcoal (10 to 30),



PLATING

Plating on Aluminum

The following formulæ for plating nickel on roughened aluminum are recommended by the Aluminum Co. of Amer-

Grease is first removed from the surface by immersion in a solution con-

Sodium Carbonate 1 to 3 oz./gal. Trisodium Phosphate 1 to 3 oz./gal. Temperature about 200° F.

The article to be plated is next rinsed in water and then preferably immersed in 5% hydrofluoric acid solution for about 15 seconds to remove the last traces of alkali and prepare for the etching solution.

The etching solution depends on the chemical composition of the metal.

Formula No. 1

For etching commercially pure aluminum use:

Nickel Chloride	36	oz.
Hydrochloric Acid (sp. gr. 1.18)		gal.
Water	1	gal.

The dipping time should be determined by actual trial. It approximates a halfminute.

No. 2

For etching aluminum alloys containing copper, manganese, and perhaps magnesium use:

Hydrochloric Acid	
(sp. gr. 1.18)	1/3 gal.
Water	% gal.
Manganous Sulphate	½ oz.
Temperature 90° F.	

The dipping time should be determined by actual trial. It approximates a halfminute.

No. 3

For etching aluminum cas	stings use:
Nitric Acid (sp. gr. 1.42)	3 fl. oz.
Hydrofluoric Acid	1
(48-52%)	1 fl. oz.
Temperature 75-80° F	

The dipping time should be determined by actual trial. It approximates a halfsolution should be lead lined and coated with the following mixture:

Beeswax	1	oz.
Paraffin	4	oz.

After etching the articles, they should be well rinsed in water, after which they may be plated in a nickel bath of formula given in Volume II.

Anodic Treatment of Aluminum

The aluminum or aluminum alloy is made the anode in a chromic or sulphurie acid solution, and 10-100 amperes per square foot is passed through for 10-20 minutes.

Formula No. 1

The chromic acid solution contains 5-15% chromic acid. The current density for this bath varies from 10 amperes per square foot to 100 amperes per square foot. The temperature of this bath is important and should be kept between 90-100° F.

Fumes of chromic acid develop as the process continues. A ventilating system should be in operation at all times as the fumes are injurious.

No. 2

The sulphuric acid method consists of anodizing the aluminum or its alloy in a solution containing 5-60% sulphuric acid by volume. The current density varies from 10 to 25 amperes per square foot. The temperature control is not as important as in the chromic acid solution.

Sulphuric acid spray is released during the process, and for this reason the bath should have a ventilating system

applied to it. After the work has been removed from the solution, it is essential to wash with water until all traces of sulphuric acid or chromic acid have been removed. For this purpose two rinses in running water for 10 minutes each will suffice.

Anodic Coating of Aluminum Formula No. 1 British Patent 427,308

The electrolyte consists of an acid to minute. The container for this etching | which a glucoside or hydrolyzed glucoside has been added. A suitable bath consists of 100 l. sulphuric acid of sp. gr. 1.220 to which is added 300 g. baptisin or 500 g. hydrolyzed barbaloin. Alternatively, 500 g. trihydroxymethylanthraquinone as obtained from the hydrolysis of frangulin may be added.

No. 2 British Patent 429.344

Caustic Soda	20 g. 1 l.
Water	1 Ĭ.
Glycerin	150 cc.

In place of the glycerin any one of the

ollowing may be used:	
Formaldehyde	75 cc.
or	
Lactose	90 g.
or	
Barbaloin	50 g.

Operate at 10-15 volts; current density 18-24 amperes per square foot at 15-25° C.

Coloring Aluminum

If anodized aluminum is placed in a solution of an organic dye, the dye unites with the coating formed on the aluminum and forms a colored lake. These colors will not wash out. Thus, by dipping anodized aluminum in a green dye solution, a green coating is obtained. In this way any desired color can be obtained.

Formation of Noncorrosive Film on Aluminum, Magnesium or Their Alloys

Japanese Patent 109,261

Aluminum, magnesium or their alloys are boiled in a solution of 25 g. of ammonium molybdate and 25 g. of ammonium tartrate per liter.

Antimony Plating

Antimony Oxide	60 g.
Hydrofluoric Acid	114 g.
Water	1000 cc.
Aloin	¼ g.
Clovel Oil	⅓ g.

The mixture should be stirred until solution of the oxide is complete. A lead vessel can be used. Vessels of these materials or of wax can be used as containers for the final plating bath. Wax vessels cannot be used in the making of the bath due to heat of the reaction. A cast antimony anode is used. This bath must be electrolyzed for several days, perhaps to eliminate impurities, before good deposits can be obtained.

A current of 0.8 ampere per sq. dm. (7.4 amperes per sq. ft.) can be used. Higher currents give less smooth deposits. Deposits can be made any thickness even 1 cm. (0.4 in.) or more. The current efficiency at the cathode is practicaly 100%.

Brass Plating

Copper Cyanide	4.2	OZ.	per	gal.
Zinc Cyanide	1.5	oz.	per	gal.
Sodium Cyanide	6.7	OZ.	per	gal.
Sodium Carbonate	4	OZ.	per	gal.
Ammonium Hv-			-	•

droxide 0.12 oz. per gal. Use brass anodes and 2-4 amperes per square foot.

Bronze Electroplating Bath British Patent 412,277

O O I-	40
Copper Cyanide	40 g.
Sodium Stannate	20 g.
Sodium Cyanide	35 g.
Caustic Soda	5 g.
Water	to make 1 l.

Brass and Bronze Solutions

Brass Solution:

Copper Cyanide	4 oz.
Zinc Cyanide	1 oz.
Sodium Cyanide	6 oz.
Sodium Carbonate	2 oz.
Water	1 gal.

Temperature 90° F. Cathode current density 2.5 to 3 amperes per sq. ft.; 2 to 3 volts. Use rolled anodes, 80% copper, 20% zinc.

Bronze Solution:

Copper Cyanide	4 oz.
Zinc Cyanide	½ oz.
Sodium Cyanide	5 oz.
Sodium Carbonate	2 oz.
Rochelle Salts	2 oz.
Water	1 gal.

Temperature 95° F. Cathode current density, 2 to 2.5 amperes per sq. ft.; 2 to 3 volts. Rolled bronze anodes, 90% copper, 10% zinc.

Cadmium Solution:

Sodium Cyanide	9 oz.
Cadmium Oxide	3 oz.
Caustic Soda	2 oz.
Water	1 gal.

Temperature 80° F. Cathode current density, 8 to 10 amperes per sq. ft.; 2 to 21/2 volts. Use iron and cadmium anodes, one iron to three cadmium. Remove cadmium anodes when solution is not in use.

Cadmium Plating Bath

Formula No. 1

Cadmium Oxide 3 oz. per gal. Sodium Cyanide 10 oz. per gal.

No. 2

Cadmium Oxide Potassium Cyanide	$39.4 \\ 128.2$	
Sodium Sulphate	50	g.
Nickel Sulphate	1	g.

Cadmium-Zinc Alloy Plating

Satisfactory deposition is possible from solutions containing 55-75 g. of zinc, 5-30 g. of cadmium, 3-6 mg. of gelatin or caffeine, and 15-20 g. of aluminum sulphate per liter, operated at 25° with pH 4 and current density 1-2 amperes per square decimeter. The cadmium content of the alloy is increased by rotating the cathode and raising the temperature and is decreased by raising the current density, increasing the acidity, and using addition agents and salts. Complex organic nitrogen addition compounds, e.g., caffeine and aloin, have a effect, retarding selective cadmium deposition and thus permitting the cadmium of the bath to be increased. Alloys containing 45-55% of zinc show most resistance to corrosion by aqueous sodium

Cadmium Plating Die Castings

Scratch brush raw die casting wet or if rough, polish first. Articles are then cadmium plated and given either a dry or wet scratch brush for desired finish. Lacquer to protect finish. Satisfactory deposits may be obtained from the following solution:

Sodium Cyanide 7 oz./gal. Cadmium Oxide 3 oz./gal. Potassium Hydroxide 2 oz./gal. Temperature 113° F. Current density 10–25 amp. per sq. ft.

Any patented brightener may be used. Strip, 10% ammonium nitrate.

Chromium Plating

The chromic acid salt to be used should consist (according to British Standard Specification) of

Chromium Trioxide (CrO₃) 99.5 % Sulphate (as Sulphuric Acid) 0.2 % Chlorides (as Chlorine) 0.05% Insoluble Matter 0.15%

and the solution made up of 250-500 g. per liter, with a density of 25 to 27° Bé. Sulphate is added in a proportion of 1/100th of the chromic acid concentration; with too high amount of sulphate, current and throwing power fall off badly. Fluoride may be substituted for sulphate, calcium fluoride 30 g./l. in a 500 g./l. solution gives good results.

The solutions should be made up very carefully; usually the bath works best when aged artificially. The tank for the solution (of glass, wood, lead-lined metal) should be arranged for heating as temperature is a critical condition; 40° C. (100° F.) is usually applied, sometimes 60° C. (140° F.) may be required, while for thick, dull deposits cold solution can be used.

Anodes are of lead or lead-antimony alloy: the latter is less affected when the bath is not operating. Current density is very important; for bright deposits on nickel 150 amperes per sq. ft., for thick deposits 300-400 amperes per sq. ft. are used. The high current density requires a particularly careful suspension of the work in the bath, thin wires as in other plating practice are out of the question; very often special jigs are used. In certain cases, where the work is rather large, auxiliary anodes of lead or iron are arranged to insure a good deposit inside a hole, recess, etc. Degreasing in trichloroethylene, polishing and nickelplating before chromium plating is desirable. Careful subsequent treatment is essential to avoid corrosive effects of eventually remaining bath solution; repeated rinsing alternately in hot and cold water, drying in an oven or hot sawdust is necessary.

Chromium Plating Bath

Chromium Oxide (Free from Sulphuric Acid) 350 g.
Potassium Fluoride 3 g.
Water 1000 cc.

Run at 18-20° C., using 3.8 to 4 volts. Chromium Solutions

Formula No. 1

Chromic Acid 33 oz.
Sulphuric Acid 0.3 oz.
Water 1 gal.

Total sulphate (from both the chromic acid and the sulphuric acid) should be 0.3 oz.

Temperature 113° F. Cathode current density 125 to 1750 amperes per sq. ft.

No. 2

Chromic Acid 55 oz.
Sulphuric Acid 0.55 oz.
Water 1 gal.

Total sulphate (from both the chromic acid and the sulphuric acid) should be 0.55 oz.

Temperature 95° F. Cathode current density 75 to 125 amperes per sq. ft.

The anodes and temperature control coils should be of 6% antimonial lead. The chromic acid tanks should be of steel, lined with 6% antimonial lead.

No. 1 is used where heavy deposits are

desired.

No. 2 is used where the deposit is for decorative purposes.

Cobalt Plating Bath British Patent 427,458

Cobalt Chloride		40-150	g.
Sodium Acid Fluoride		10 - 40	ğ.
Ammonium Chloride		15 - 60	
Cobalt Basic Acetate		15-60	ğ.
Water	to	make 1	Ĩ.

Copper Solutions

Cyanide Copper Solution	No.	1
Copper Cyanide	$3\frac{1}{2}$	oz.
Sodium Cyanide	$4\frac{1}{2}$	oz.
Carbonate of Soda	2	oz.
Hyposulphite of Soda	1/64	oz.
Water	1	gal.
No 2		_

Copper Carbonate	5 oz.
Sodium Cyanide	10 oz.
Hyposulphite of Soda	½4 oz.
Water	1 gal.

Either solution should be operated at 100° F. to 110° F. Cathode current density 4 to 6 amperes per sq. ft.; 1½ to 2 volts. Use rolled copper anodes.

Acid Copper Solution

Copper Sulphate	28 oz.	
Sulphuric Acid	3 to 5 fl. oz	
Water	1 gal.	

Temperature 75° F. Cathode current density for still solution 10 to 15 amperes per sq. ft.; ¾ to 1 volt. Agitation of the cathode or of the solution allows the use of higher current density. Use rolled copper anodes.

Coppering by Immersion

Copper Sulphate	1	to	2	oz.
Sulphuric Acid	1/2	to	1	OZ.
Water			1	gal.

Where only a very thin film of copper is desired the above solution will give good results.

Acid Copper Plating

Cupric Sulphate 27 oz. per gal. Sulphuric Acid 7 oz. per gal.

Use brass anodes and a current density of 20-40 amperes per sq. ft.

Cyanide Copper Plating

T, amide cop	POL I	1 CL L I I	us .		
Copper Cyanide	4	oz.	per	gal.	
Sodium Cyanide	5.5	OZ.	per	gal.	
Sodium Carbonate				gal.	
Use at 35° C. at 13	amp	eres	per	sa. f	t

Blue Dip (for Plating Copper and Brass Articles)

Bichloride of Mercury	1/2	oz.
Sodium Cyanide	6	OZ.
Ammonium Chloride	1	oz.
Water	1	gal.

Fluoride Bath

	g.		
Antimony Oxide			
(Commercial)	60	8	oz.
Hydrofluoric Acid			02.
(Commercial,			
48%)	114	15.3	oz.
Water	1000	1.6	gal.
Aloin	0.25	0.033	oz.
Clove Oil		0.0016	oz.
	3.012		~~

The last two constituents, the so-called addition agents, are used up during the plating; hence, they must be added regularly to the bath. The quantities given above are sufficient for about 12 hours of operation.

Immersion Gold Solution

Fulminate of Gold	4	dwt.
Yellow Prussiate Potash	24	oz.
Carbonate of Soda	12	oz.
Caustic Soda	1/4	oz.
Water	1	gal.

Solution should be boiled in a cast iron tank for an hour and allowed to cool to 180° F. before using.

Salt Water Gold

Yellow Prussiate of Potasl	64 oz.
Sodium Phosphate	32 oz.
Sodium Carbonate	16 oz.
Sodium Sulphite	8 oz.
Gold as Fulminate	12 dwt.
Water	4 gal.
Solution is boiled for one	hour, ther

diluted with water to make 4 gal. of solution. The solution is placed in a porous pot which is put in a tank that contains a saturated solution of sodium chloride heated to 190° F.

Green Gold

Metallic Gold as Fulmi	_	
nate or Cyanide		4 dwt.
Silver Cyanide		¼ dwt.
Sodium Cyanide		2 oz.
Carbonate of Soda		2 oz.
Water		1 gal.
Temperature 105° F.	: 2	volts: 1

Temperature 105° F.; 2 volts; 18 karat green gold anodes.

Rose Gold

Yellow Prussiate of Potash	4	OZ.
Potassium Carbonate	4	0 Z .
Sodium Cyanide	1/4	OZ.
Gold as Fulminate	10	dwt.
Water	1	gal.

Temperature 175° F.; 6 volts. If a red color is desired, add small quantity of copper carbonate.

Coating Iron with Lead and Tin

Iron and steel can be coated electrolytically after pickling with sulphuric acid, in a bath of tin borofluoride, lead borofluoride and borofluoric acid with acid-proof layers of a lead-tin alloy which are so elastic that the metals can still be worked mechanically; the temperature must, however, not rise above 150 to 200° C., as otherwise the coatings would melt. The deposits are made at a current density of 0.5-3.0 amperes per sq. dm.

Electrolytic Burnishing of Iron

Oxidize anodically in 20 to 40% caustic soda at 1 to 6 amperes per sq. dm. at 1 to 2 volts at 60-70° C.

Thin Deposits of Iron

Dissolve 16 oz. of ammonium chloride in each gallon of water. Connect up tank, same as for plating, using cold rolled iron for anodes. On the cathode rod suspend some old plating racks or other work, and work solution with highest current density obtainable. After 4 or 5 hours of work of the solution, there will be enough iron dissolved from the anodes and the solution will produce a deposit of iron. Operate solution at 80° F.; 1.5 to 2 amperes per sq. ft.; 1 volt.

Tron Solution

Ferrous Chloride	40	OZ.
Calcium Chloride	20	oz.
Water	1	gal.
m 4 0000 TI		7

Temperature 200° F.; current density 40 to 50 amperes per sq. ft.; 2 to 2½ volts; pH 1.5 to 2; pure iron anodes.

This bath is used to produce heavy deposits of iron.

Preparing High-Speed Steels for Plating

In order to secure good adhesion of electro-deposits to high-speed steel it is treated anodically at 2.7 amperes per sq. dm. in a bath containing 115 g. of caustic soda and 15 g. of citric acid per liter until gas evolution is uniform over the whole surface, then rinsed with water, dipped momentarily in 6-12N-hydrochloric acid and finally washed with water.

Electrodeposition of Lead

Fifty grams of lead perchlorate and 10 g. perchloric acid in 1 liter electrolyte and a current density of 0.25-0.50 amperes per square decimeter are recommended for the preparation of pure 0.1 mm. deposits of lead of good texture. Agitation of the bath permits a higher current density and thicker deposits. Addition of 0.2-0.4 g. peptone and moderate agitation improve the deposit and allow a current density of 1 ampere per square decimeter. Higher current densities up to 2 amperes per square decimeter require constant and efficient stirring and heating up to 60° C. permits 3-4 amperes per square decimeter. For technical purposes 1 ampere per square decimeter is recommended.

Lead Solutions

Lead Carbonate		20	oz.
Hydrofluoric Acid (50%)	32	oz.
Boric Acid	•	14	oz.
Glue		0.025	02

Place the hydrofluoric acid in a leadlined tank and add the boric acid with constant stirring. When the boric acid is completely dissolved, the solution is allowed to stand until cool, when the lead carbonate is added in the form of a paste with water. The solution is allowed to settle in the plating tank. The solution is then diluted to the proper volume with water and the glue added after dissolving the same in warm water. Mechanical agitation of the solution is essential.

A cathode current density of 10 to 20 amperes per sq. ft., 3 to 4 volts, and lead anodes are employed.

Thin Deposits of Lead

Carbonate of L	കെറ്	9	oz.	
Caustic Soda	cau	-	OZ.	
		-		
Water			ga	ı.
Lead anodes.	Temperature	17:	ĭ٥	F

3 to 4 volts.

Coating Magnesium and Its Alloys French Patent 766,685

Magnesium or an alloy thereof is coated by introducing it into a rotating drum along with an alloy of zinc (25) and cadmium (75 parts) and some galvanized iron turnings. The drum is heated to about 290° C., when the alloy becomes pasty, and is rotated for about 3 minutes.

Commercial Nickel Plating

The three principal methods of nickel plating, i.e., ordinary plating in the stationary bath, rapid plating and barrel plating are discussed and compared as to their respective economic advantages. In all methods it is necessary that new nickel sulphate be continuously formed at the anode and that the deposit be fine in grain. The deposit must permit of mechanical working without injury. The deposit if chromium plated must not peel. The composition of an ordinary stationary bath consists of 75 g. nickel ammonium sulphate in one liter water with a pH of about 5.8; increasing the latter to 6.4 increases, reducing it to 4.6 decreases the throwing power of the bath. Specific gravity is 6-7° Bé., the current density 0.3 ampere per square decimeter, voltage 3.5, temperature 18° C. A thickness of 0.025 mm. is obtained in 7 hours. A rapid plating bath must work at 50°, the grain of such deposit is the finer, the better the electrical conductivity of the bath. The compositions used are: 240 g. nickel sulphate, 30 g. boric acid. 270 g. meket surpnate, 30 g. boric acid, 19 g. potassium chloride in 1 liter water; or 240 g. nickel sulphate, 120 g. magnesium sulphate, 30 g. boric acid; or 240 g. nickel sulphate, 30 g. boric acid; acid, 150 g. magnesium archetet 10 g. magnesium archetetetes 10 g. supplied acid, 10 g. s 150 g. magnesium sulphate, 10 g. sodium chloride, 50 g. sodium sulphate, 0.1 g. sodium fluoride in 1 liter water. The current density must be adapted to the kind of ware to be plated. Pure nickel anodes do not dissolve as easily as 98% nickel anodes. If the deposition velocity is too high, an excess of oxygen is formed at the anode, passivates it and finally nickel bisulphate and peroxide are formed without nickel going into solution. Plating in the barrel requires a pH of not less than 6.6, at 8-12 volts, time usually 2 hours, bath temperature 35-50°.

Nickel Solutions

Nickel Solution for Brass, Copper, and Cold Rolled Steel

A nickel solution that has been used with good results on brass, copper and cold rolled steels is made as follows:

Formula No. 1

Double Nickel Salts	8 oz.
Single Nickel Salts	4 oz.
Boric Acid	2 oz.
Sodium Chloride	2 oz.
Water	1 gal.

Solution to be operated at 80° F.; 2 to 2½ volts; 6 to 8 amperes per sq. ft. and a pH of 5.8.

For solutions that are operated at a higher temperature and a correspondingly higher current density, use:

No. 2

Double Nickel Salts	8 oz.
Single Nickel Salts	8 oz.
Sodium Chloride	3 oz.
Boric Acid	3 oz.
Water	1 gal.

Temperature 110° F.; 2½ to 3 volts; 20 amperes per sq. ft., and a pH of 6; depolarized nickel anodes 99% plus. Replenish by the addition of single nickel salts.

Low pH Solution for Heavy Deposits of Nickel

No. 3

Single Nickel Salts	32 oz.
Sodium Chloride	6 oz.
Boric Acid	4 oz.
Water	1 gal.

Nickel Strip

Sulphuric Acid	4	oz.
Water	1	oz.

Temperature 80° F.; lead cathodes; 6 volts. If 3 or 4 oz. of copper sulphate per gallon are dissolved in the water before adding to the acid, the strip will not attack the base metal so readily.

Nickel Brighteners

Bright deposits of nickel are obtained from No. 1 formula above by the use of cadmium chloride or one of the prepared brighteners that are on the market. The pitting of nickel deposits is eliminated by adding hydrogen peroxide to the bath. Use from 1 to 5 cc. of 100 volumes peroxide to each gallon depending upon the severity of the pitting.

Nickel Plating

The nickel content of the bath is about 40-50 g. per liter; current density 0.3-0.4 amperes per square decimeter while for rapid plating methods 1-3 amperes per square decimeter are employed. The bath is stirred and the pieces are moved to avoid streaks on the deposit, pH is 5.8-6.2. For rapid nickel plating the following bath is recommended: pure nickel sulphate 22.5 kg., pure ammonium sulphate 2.0 kg., pure nickel chloride 0.5 kg., pure sodium perborate 0.5 kg., water 100 liters 35-40° C., voltage 2.75-3.5.

Hydrogen Poor Nickel Plating

Nickel sulphate 80 g., nickel fluoride 8 g., sodium chloride 1 g., sodium sulphate 0.5 g., sodium nitrate 0.02 g., sulphosodium-phenolate 0.12 g., sodium citrate 2 g., boric acid 6 g., zircon-ammonium fluoride 0.2 g., all in 1 liter water. The ammonium fluoride binds the hydrogen and the deposits adhere well to the base. The voltage employed with this bath is 2 volts.

White Nickel Plating Formula No. 1 (Low Metal Bath)

Nickel Sulphate
Ammonium Chloride
Boric Acid
pH = 5.4

12 oz. per gal.
2 oz. per gal.

Use at room temperature with nickel anodes, and 10-20 amperes per sq. ft.

No. 2 (High Metal Bath)

Nickel Sulphate
Nickel Chloride
Boric Acid
pH = 5.3

34 oz. per gal.
4 oz. per gal.

Use nickel anodes and a current density of 15-45 amperes per sq. ft. with a temperature of 50-60° C.

Nickel Bath for Die Castings

Temperature, 20-30° C.; current density = 15-30 amperes per sq. ft.

Depositing Nickel on Rough Steel If a smooth deposit is required over rough steel, instead of buffing down the steel, it is possible to pickle the steel in an acid until all the scale is removed and then depositing a heavy coat of copper, using an acid sulphate bath for this purpose. The heavy coat of copper is then buffed until it is smooth. The coat can now be finished in any way desirable. It is much cheaper to buff copper than steel.

Black Nickel Plating

Nickel Ammonium Sulphate 60 g. per l. Zinc Sulphate 14 g. per l. Sodium Sulphocyanate 14 g. per l. pH \Longrightarrow 5.8-6.0

Gray Nickel Plating

Nickel Ammonium
Sulphate
Sodium Sulphocyanate
pH = 5.4

Sodium Sulpho14 g. per l.

Plating Zinc with Nickel

(1) Strike for 5-10 minutes in any suitable cold nickel solution. The following formula is suggested:

Nickel Sulphate 15 oz. per gal. Anhydrous Sodium

Sulphate 15-18 oz. per gal. Ammonium Chloride 2-3 oz. per gal. Boric Acid 2 oz. per gal. Temperature 78-85° F. pH = 4.9-5.4 (electrometric)*

Current density 24-30 amp. per sq. ft. (2) Rinse thoroughly in cold water.

(3) Transfer without drying to the following solution:

Nickel Sulphate 20 oz. per gal. Ammonium Chloride 4 oz. per gal. Boric, Acid 2 oz. per gal. Temperature 105-115° F. pH = 5.0-5.3 (electrometric)

Current density 40-80 amp. per sq. ft.

* May be increased to as high as 30 ounces
per gallon for intricate shapes.

Solvent Cleaning of Zinc

Grease and oil may be removed from zinc and zinc alloy castings by the use of trichloroethylene, carbon tetrachloride, xylol, ethyl acetate, etc. These solvents are most effective when used in apparatus involving vapor rinsing. However, these solvents do not remove oxide films and zinc salts and hence where parts are to be electroplated, the metal should subsequently be submitted to an

acid dip which serves the additional purpose of roughening the surface to provide good adhesion of the finish coating. The following solutions have been used in zinc alloy die castings:

(1) Phosphoric acid etch-treat for 30 seconds in 3% solution of phosphoric acid (85% H₂PO₄ grade, specific gravity

1.74) rinse and dry.

(2) Hydrochloric acid etch-treat for 30 seconds in a 10% solution of hydrochloric acid (35 to 37% HCl grade, specific gravity 1.18-1.19) rinse and dry.

(3) Hydrofluoric acid etch-treat for 30 seconds in a 1% solution of hydrofluoric acid solution (48% HF grade) rinse and dry.

Plating of Zinc

Considering nickel and nickel-chromium plated coatings on zine and zine alloy castings, a minimum thickness of coating of 0.0003 in. at the thinnest point is necessary to give any satisfaction in outdoor service. Completely satisfactory quality will not be obtained consistently with coatings of less than 0.001 in. average thickness.

Nickel Plating Solutions Formula No. 1

Nickel Sulphate 10 oz. per gal. Anhydrous Sodium

10-15 oz. per gal. Sulphate Ammonium Chloride 2-3 oz. per gal. 2 oz. per gal. Boric Acid

Operating details for this solution fol-

pH-This should be held between 5.3 and 5.7 electrometric or 5.8-6.2 colorimetric. The anode area should be controlled to minimize pH changes. pH should be checked daily and adjustments made by the addition of ammonium hydroxide or sulphuric acid as needed. Under best operating conditions this solution will tend slowly to become alkaline.

Temperature—For use in applying nickel directly on zinc this solution should be kept at or preferably slightly above room temperature (70-80° F.). If the temperature falls below 70° F. the deposits will be hard and brittle showing cracks. Temperature above 80° F. will tend to cause the formation of black streaks in recesses

Nickel Content-The prescribed nickel sulphate content corresponds to about 2 oz. per gallon of nickel calculated as metal. No harm will result if this increases somewhat in use.

Sodium Sulphate Content-The amount of sodium sulphate present in the solution should be regulated to suit the complexity of the articles to be plated. Simple shapes may require not more than 10 oz. per gallon of sodium sulphate. More complicated shapes may require the presence of 15 oz. per gallon or more. Some commercial platers add as high as 30 oz. per gallon. In general, the sodium sulphate content should be the lowest possible for the articles being plated.

Current Density-When made up according to the formula given, the bath should be operated at between 12 and 20 amperes per sq. ft. The maximum current density will be determined by the tendency for the deposits to burn. In the presence of very high sodium sul-phate concentrations, burning may develop at current densities lower than 20 amperes per sq. ft. If streaking occurs at the maximum current density, purifica tion of the solution may be necessary.

Agitation—Agitation reduces porosity and permits the use of somewhat higher current densities. With certain shapes, agitation will be found absolutely necessary for successful plating.

Pitting-Like all other nickel solutions this bath will at times develop a ten-dency towards pitting. This is usually an indication that foreign matter is present. A temporary cure can be effected by adding hydrogen peroxide or sodium perborate to the solution. Permanent freedom from pitting can only be obtained by continuous filtration and scrupulous care in avoiding the presence of foreign material in the solution. Pitting may on occasion develop from faulty cleaning.

No. 2

Nickel Sulphate 15 oz. per gal. Anhydrous Sodium

Sulphate 15 oz. per gal. Ammonium Chloride 3 oz. per gal. Boric Acid 2 oz. per gal.

Operating details for this solution are given below:

pH-Should be kept between 4.9 and 5.4 electrometric or 5.4-5.9 colorimetric by means of additions of sodium hydroxide or hydrochloric acid. Ammonium hydroxide and sulphuric acid should not be used as the solution is nearly saturated with respect to nickel ammonium sulphate.

Temperature-The more concentrated solution permits the use of somewhat higher current densities which in turn permit the use of higher temperatures of operation which may be reflected in slightly softer deposits. The minimum safe temperature is 75° F. and the maximum is 87° F.

Nickel Content—Corresponds to about 3 oz. per gallon calculated as nickel metal. Any large increase in nickel content may result in crystallization of double nickel salts from solution.

Sodium Sulphate Content—Should be regulated as for the 2-oz. (nickel content) solution. In general, somewhat higher sodium sulphate contents will be required in the present case.

Current Density—This more concentrated solution permits the use of higher current densities, the range in the present case lying between 24 and 36 amperes per sq. ft.

Agitation-Pitting—The considerations mentioned under Formula No. 1 above hold in the present case.

No. 3

Nickel Sulphate 20 oz. per gal. Ammonium Chloride 4 oz. per gal. Boric Acid 2 oz. per gal.

Operating details for this solution are given below:

pH—The pH of this solution should be held between 5.0 and 5.3 electrometric or 5.5-5.8 colorimetric. Higher pH will tause cracking and peeling while lower pH will tend to increase the attack of the solution on exposed portions of the base.

Temperature—Should be between 105 and 115° F. (40-45° C.). Lower temperatures will not permit the deposition of soft nickel. Higher temperatures, while allowable, tend to cause excessive loss of water by evaporation.

Current Density—The current density should under no circumstances fall below 40 amperes per sq. ft. and preferably should be maintained at 60 amperes per sq. ft. or higher. Not only does the speed of production fall off at the lower current densities but contamination of the solution becomes more serious. These current densities are similar to those required for chromium plating and suitable generator capacity should be available.

Agitation—Agitation will tend to reduce pitting and porosity.

Pitting—Like most warm solutions, new baths of this composition may develop an exaggerated type of pitting. This condition can be readily overcome by additions of hydrogen peroxide. Sodium perborate should never be used for the reasons given below.

Sodium Salts—Sodium salts should not be permitted to enter this solution. When the solution is pure, very high current densities can be employed without burning. The presence of sodium salts very definitely restricts the operation to low current densities which not only do not utilize the full production capacity of the solution but also permit excessive zinc pickup. For these reasons the rinsing between nickel tanks should be thorough, sodium perborate should not be used to prevent pitting, and additions of alkali to raise pH should be made with ammonium hydroxide rather than sodium hydroxide.

Nickel Plating Methods

Three methods of applying adequate nickel coatings to zinc and zinc alloy castings have been found successful.

Multiple Nickel

This method consists essentially of depositing on the zinc articles, from either Formula No. 1 or No. 2 above, a coating of nickel 0.0001 in. to 0.0002 in. thick following which the articles are thoroughly rinsed in cold water and placed in a warm nickel solution (Formula No. 3) for completion of the plating to the required thickness.

The strike coating must be adequate to protect the zinc base from the action of the subsequently used warm solution. For simple shapes a 5-minute deposit at 25 amperes per sq. ft. may be sufficient. More complicated shapes will need 10 minutes at this current density.

Rinsing—In the interval between the two nickel tanks the articles should not be allowed to dry. If drying does occur poor adhesion of the second coat will develop. The use of cold water in the rinse will minimize the danger of this happening.

Copper-Nickel

While the system of plating nickel direct has a great many advantages, good results have also been obtained commercially by plating with copper-nickel deposits totalling 0.001 in. in thickness.

In this system of plating, the work is cleaned thoroughly, a coating of copper is applied to a thickness of 0.0005 in. from a copper-cyanide solution, followed, after rinsing, by the application of 0.0005 in. of nickel in a warm nickel solution (Formula No. 3).

The copper cyanide solution may be any one of those commonly used. A typical formula follows:

Sodium Cyanide	4-6 oz./gal.	(30-45	g. per l.
Copper Cyanide	4 oz./gal.	(30	g. per l.
Sodium Bicarbonate			g. per l.
Sodium Bisulphate	¼ oz./gal.	(1.87	g. per 1.)

The solution should be used at 70–113° F. (21–45° C.) with a current density of 10–15 amperes per sq. ft.

The copper-nickel system of plating is adapted to the production of heavy deposits. Its use is not advocated for coatings less than 0.0005 in. in thickness. The copper layer should be at least 0.0002 in. thick in order to avoid complete absorption by the zinc base and to provide protection of the zinc base from attack by the warm nickel solution. The copper layer fills the same role here as the primary or strike nickel deposit in the multiple nickel system of plating.

The nickel deposit must be at least 0.0003 in. thick for outdoor use. Thinner deposits will readily permit the seepage through pores of copper salts which will stain the surface with an unsightly brown

film.

Nickel-Copper-Nickel

When coatings ranging from 0.00075 in upward are desired, multiple coatings are necessary to avoid cracking. Multiple nickel coatings have been described above. The system nickel-copper-nickel has also been used successfully.

Clean the articles thoroughly as described under "Cleaning of Zinc and

Zinc Alloys."

Plate 0.0002 in. of nickel in either cold solution described in Formulas No. 1 and No. 2.

Plate 0.0004 in. of copper from an acid-copper solution.

Color copper, coat, and clean.

Plate 0.0004 in. of nickel from any warm nickel solution such as described in Formula No. 3 above.

The buffing operation is not essential if the two primary coats are sufficiently smooth to make coloring of the final nickel readily accomplished.

The acid copper solution may be of any accepted composition. The follow-

ing formula is typical:

Copper Sulphate 24 oz./gal. Sulphuric Acid 6–8 oz./gal.

This solution is used at room temperature to 113° F. (45° C.) with a current density of 10-50 amperes per sq. ft. Animal glue may be used as a brightener in amounts of ½ oz. per gal. (0.9 g. per l.).

Bright Nickel Plating on Zinc

A bright nickel deposit which requires no buffing or coloring can be produced in the sulphate type of solution by the addition of ½00 of an oz. per gal. of cadmium sulphate. A small amount of cadmium sulphate may be added from time to time to maintain the cadmium metal content in use.

The deposits produced are very bright and smooth but somewhat brittle and should not be deformed or bent. Chromium should not be deposited over such coatings as the additional stress will

crack and peel the nickel.

Bright nickel deposits of this type tend to be brittle and are suitable only for use in thin form for indoor application.

Black Nickel Plating on Zinc

A bright, black, adherent coating can be obtained on zinc by a 2-minute plating in the following solution at 113° F. (45° C.).

Nickel Ammonium
Sulphate 8 oz./gal.
Zinc Sulphate 1 oz./gal.
Sodium Sulphocyanate 2 oz./gal.
Current Density 1-2 amp./sq. ft.

Chromium Plating * on Zinc

Chromium may be applied either as a thin finish coating over nickel or as a heavy protective coating directly on zinc from the following solutions:

 $\begin{array}{c} \text{Chromium Oxide (CrO}_3) & 33 & \text{oz./gal.} \\ \text{Sulphuric Acid } (\text{H}_2\text{SO}_4) & 0.3 & \text{oz./gal.} \\ \text{Or} \\ \text{Chromium Oxide (CrO}_3) & 33 & \text{oz./gal.} \\ \text{Chromium Sulphate} \\ (\text{Cr}_2(\text{SO}_4)_3) & 0.44 & \text{oz./gal.} \\ \end{array}$

For finish plating this should be used at 113° F. (45° C.) with lead anodes and at a current density of 75–150 amp. per sq. ft. A 3-6 minute deposit should be sufficient.

For heavy deposits applied directly on zinc these solutions may be used with the conditions of operations stated. The

* No consideration has been given to the patent situation involving chromium solutions which must be taken into account by the plater,

work should be plated for 20-25 minutes to insure reasonable thickness of coating. The deposits obtained will not be bright but will have a luster ranging from milky to frosty depending upon conditions. The explanation for the failure to obtain bright deposits apparently lies in the fact that these solutions etch the surface of the zinc slightly before deposition occurs to protect it. The deposits can, if only milky, be readily buffed to a bright luster.

Somewhat better protection and ease of buffing will be obtained with chromium deposits applied directly on the zinc from these solutions at room temperature with a current density of 50–125 amp. per sq. ft. The deposits will be dull gray in appearance but can be readily buffed or brushed to a high luster. The work should be plated for 20–25 minutes to insure a good protective plate.

Cadmium Plating * on Zinc

Recent practice to improve the surface appearance of zinc alloy die castings such as carburetors, etc., which do not require a fine finish is to cadmium plate them directly without buffing. Satisfactory deposits may be obtained from any of the numerous types of solution in use. A typical formula is:

Sodium Cyanide	7 oz./gal.
Cadmium Oxide	3 oz./gal.
Caustic Potash	2 oz./gal.

This solution should be used at room temperature to 133° F. (45° C.) with a current density of 10-25 amp. per sq. ft. Almost any of the patented brighteners will give satisfactory results.

* No consideration has been given here to the patent situation involving cadmium plating which must be taken into account by the plater.

Stripping Methods

Nickel-Chromium

Chromium and nickel may be removed by making the work anode in concentrated sulphuric acid to which a small quantity of commercial glycerin is added. Zinc is only slowly attacked by the concentrated acid but as the solution absorbs moisture from the air this attack will increase to the point where pitting of the zinc starts and the solution demands attention. The excess moisture may be removed by boiling the solution until heavy white fumes appear.

Nickel Coatings

Immerse in the following cold solu-

Water	1	part
Sulphuric Acid	2	parts
Nitric Acid	2	parts
Hydrochloric Acid	1/16	part

Prepare by adding the sulphuric and nitric acids to water and, after allowing the solution to cool, adding the hydrochloric acid.

Non-Electric Nickel Plating Compound Formula No. 1

_ 077774740 _ 1.01	
Nickel Ammonium Phosphat	e 5 oz.
Nickel Sulphate	3 oz.
Cream of Tartar	2 oz.
Tin Chloride	2 oz.
Ammonium Chloride	1 oz.
Codium Chloride	1 oz.
Copper Powder	2 oz.
Challe Dowdon (Whiting on	

Chalk Powder (Whiting or Precipitated Carbonate) 4-5 oz. Water until pasty

No. 2

Nickel Ammonium Sulphate Nickel Sulphate Cream of Tartar Tin Chloride	25 g. 15 g. 10 g. 10 g.
Ammonium Chloride	5 g.
Salt	3 g.
Whiting	20 g.
Metallic Copper, Powder	10 g.
Water until	pasty

Rhenium Plating

Rhenium, with an atomic weight of 186.3, is a very heavy metal. It is both ductile and malleable, and has a brinell hardness of 250. It is quite soluble in nitric acid but insoluble in hydrochloric acid. Therefore it should find wide use for plating on jewelry, as the hydrochloric acid released in perspiration will not affect the deposit.

Bath 1

Potassium Perrhenate 11 g. per l. Sulphuric Acid 9.3 g. per l. Temperature, 25°-45° C. (77°-113° F.) Current Density, 90-110 amp. per sq. ft.

Rath 2

			~					
Perrhenic	Acid			2	0 :	ø.	per	1.
Sulphuric	Acid				5	ø.	per	1.
Temperati	ire, 25°	°-30°	C.	(77°	8€	3°	F.)	
Current I	ensity,	90-14	40	amp.	pe	r	sq. f	ŀt.

Bath 3

Dissolve 8 g. of rhenium in concentrated nitric acid. Add 4 cc. of concen-

trated sulphuric acid, and boil until sulphur trioxide fumes are evolved. Dilute to one liter, and add enough sulphuric acid until 6 g. per l. is obtained. This solution may be used at 20°-60° C. with 50-100 amp. per sq. ft. using platinum as an insoluble anode, or rhenium as an anode. The metal deposits as a smooth shiny adhering deposit. The plating time can be 10-60 minutes.

Rhenium Nickel Plating

Potassium Perrhenate
Nickel Sulphate
Sulphuric Acid
Temperature, 25°-50° C.

11 g. per l.
6 g. per l.
9.3 g. per l.

Current Density, 50-60 amp. per sq. ft.

The alloy of nickel rhenium obtained

The alloy of nickel rhenium obtained from the above solution is somewhat lighter in color than pure rhenium.

Rhodium Plating

Five g. of rhodium chloride in 1 l. water are boiled with 40 g. of sodium nitrite until light yellow; 3 g. of sodium carbonate are added to remove traces of bismuth and the solution is filtered. After cooling 50 cc. of saturated aqueous ammonium chloride are added and precipitated ammonium rhodinitrite is collected and washed with cold water. 8.52 g. are heated to fuming with 33 cc. of concentrated sulphuric acid cooled, and diluted to 1 l. Deposition is best effected at 40° C. with platinum electrodes using a current density of 5 amp. per sq. ft. Cathode current efficiency is about 45%.

Rhodium Plating Silver Canadian Patent 343,808

Five g. of rhodium ammonium nitrate is dissolved in 1 l. of boiling water containing 20 cc. of sulphuric acid, and after the reaction is completed 100 g. sodium nitrate and 20 g. ammonium nitrate are added. The mixture is evaporated to dryness and the residue dissolved in 1 l. of water to form an electrolyte for plating silver. Deposition is preferably conducted at 80-100° F. with a current of 20-50 amp. per sq. ft. of cathode surface and an inert anode, such as carbon or platinum. The plated silver resists tarnishing.

Non-Poisonous Silver Plating

Tion T ornorroup	 	
Silver Nitrate	25-30 g.	
Thiourea	60-70 g.	
Water	1 1.	

Use 0.2 amp. per sq. dm. at 30-35° C. at 1½ volts

Silver Dip

Silver Chloride 1½ oz./gal. Sodium Cyanide 2½ oz./gal.

In order to apply this procedure to headlight reflectors it is necessary to remove any nickel plate, then polish and clean before dipping. The film of silver so produced is very thin and will have a short useful life.

Improving Silver Finish

There is no bright dip for silver in the same way as a dip for brass or copper. The surface of the parts in question can be improved by making them anodes in a solution containing 8 oz./gal. of sodium cyanide and 8 oz./gal. of sodium ferrocyanide. Use 10-15 amp./sq. ft. and about 6 volts pressure. Keep work well agitated.

Non-Poisonous Silver Plating

Citric Acid	60	g.
Sodium Iodide	520	

Use a silver anode with current density of 1-1.8 amp. per sq. dm.

Silver Plating Stainless Steel

In silver plating stainless steel it is essential to etch slightly the surface with an acid pickle. This is done to obtain a metallic surface that the subsequent electro-deposit of silver will adhere to.

A pickle made up of 10%-15% sulphuric acid, either electrolytic or still, at a temperature of 150°-160° F., will work satisfactorily.

A silver plating bath of the following composition can be used:

Silver Cyanide
Sodium Cyanide
Free Cyanide
Water

4 troy oz./gal.
5 oz./gal.
4 oz./gal.
1 gal.

Non-Electric Silver Plating Compound

Silver Nitrate	6	oz.
Ammonium Chloride	6	oz.
Sodium Thiosulphate	10	oz.
Calcium Carbonate or	Chalk 10	oz.
Water	until pa	sty

Brightener for Silver Solution

Silver Solution	1	qt.
Sodium Cyanide	8	oz.
Carbon Bisulphide	e 1	oz.
Ether	1	OZ.

To prepare the brightener place the carbon bisulphide and ether in a quart

bottle and shake thoroughly. Dissolve the cyanide in the silver solution and fill bottle. Shake bottle from time to time until the carbon bisulphide is thoroughly dissolved and then filter. One ounce of this stock solution should be sufficient for an addition to each 50 gal. of the regular plating solution. Care must be taken to avoid an excess.

Silver Strips

Formula No. 1

Codina	Cyanide	19	oz.
Caustic	Soda	2	oz.
Water		1	gal.

Reverse current with cold rolled steel as cathodes. Voltage 6 to 8. Agitate the work for a cleaner job.

No. 2

Sulphuric Acid	5	gal.
Nitric Acid	1	gal.

Place crock that contains the strip in a hot water container. If all water is kept from the strip, brass or copper work will be attacked only slightly.

Removing Fire Scale from Silver

Nitric Acid 2 oz. Water 1 oz.

Use hot and agitate work.

Removing Fire Scale by Reverse Current Sodium Cyanide

Water 1 gal. Use hot and agitate work. Lead anodes; 4-6 volts.

Bright Dip

Sulphuric Acid	2 gal.
Nitric Acid	1 gal.
Water	1 qt.

Add 1 oz. of muriatic acid for 5 gal. of above.

It is necessary to add water only when a new bright dip is made. Dip must be operated cold.

Matt Dip

Sulphuric Acid	1 gal.
Nitrie Acid	1 gal.
Zinc Oxide	2 lb.

Operate hot and keep out all water and chlorides. If the matt is coarse, add sulphuric; if too fine nitric acid.

Gold Solutions

Cyanide Solution

•		
Metallic Gold as fulminate		
or Cyanide	5	dwt.
Sodium Cyanide	2	oz.
Sodium Phosphate	1	oz.
Water	1	gal.

Temperature 130-160° F.: 1 volt; 24 kt. gold anodes.

Chloride Solution

Gold Chloride	6	OZ.
Hydrochloric Acid	10	oz.
Water	1	gal.

Room Temperature; 2-3 volts.

In preparing the solution dissolve the gold chloride in dilute hydrochloric acid before adding it to the solution.

Silver Solution Formula No. 1

Silver Cyanide $3\frac{1}{2}$ oz. Sodium Cyanide oz. Sodium Carbonate 2 oz. Water gal.

No. 2 Silver Cyanide 3½ oz. Sodium Cyanide oz. Sodium Carbonate

oz.

Water gal. Either of the two solutions will give good results if operated at a temperature of 75° F. with a cathode current density of 4 or 5 amp. per sq. ft.; 3/4 to 1 volt. Formula No. 1 is generally used, but the

Silver Strike

deposit of No. 2 is whiter.

DIIVCI	CULIE		
Silver Cyanide		1/2	oz.
Sodium Cyanide		8	oz.
Water		1	gal.

Use steel or carbon anodes; 6 volts. Black or Gun Metal Finish on Steel

A black or gun metal finish may be obtained on steel articles by heating them in a retort with a small amount of charred bone and heated to 700°-800° F. After articles are thoroughly oxidized temperature is dropped to 650° F. and a mixture of bone and bone oil is added. Several hours are required to produce finish. Articles after coming from retort are rolled in oily granulated cork until uniform black finish is secured.

The following solution will give to aluminum a uniform black color:

Water Potassium Permanganate 5-10 g. Nitric Acid 28° Bé. 2– 4 cc. Copper Nitrate 20–25 g. Temperature, 80° C.

Time to obtain deep black, 20-30 minutes.

Tantulum Plating U. S. Patent 1,933,319

The electrolyte is a fused mixture of Potassium Chloride 300 g. Potassium Fluoride 120 g. Potassium Tantulum Fluoride 100 g. Tantulum Oxide 25 g. in a graphite crucible at 750° C. This bath gives a bright plate on iron or nickel at 1 to 10 amp. per sq. dm.

Tin-Plating from An Alkaline Bath

Tin-plating of copper, brass, zine, lead, hard lead, iron, steel and aluminum can best be carried out at 0.15–0.5 volt in alkaline aqueous stannous chloride, or in alkaline aqueous sodium stannate plus sodium chloride, with 0.12–0.2 g. of gelatin per l. A tin anode (anode current density 0.45–1.6 amp. per sq. dm.) can be used. A cathode current density is 0.2–1.5 amp. per sq. dm. The maximum and minimum concentrations of the bath are 50 g. of tin salt for 2 molecules of sodium hydroxide and 12 g. for 1 molecule respectively.

Non-Poisonous Tin Bath

An alkaline tin bath without cyanides to be used at 50-60° C. is composed of sodium stannate 7.5 kg., sodium acetate 1.25 kg., sodium hydroxide 1.25 kg., starch 70 g., water 100 l. Anodes are partly of tin, partly of iron. The bath can be used for electrical tinning of kitchen utensils.

Tin Solution

Sodium Stannate	12 oz.
Caustic Soda	1 oz.
Sodium Acetate	2 oz.
Hydrogen Peroxide	1/12 oz.
(25 Volume) or	/
Sodium Perborate	½ oz.
Water	1 gal.

The solution is operated at a temperature of 140-160° F.; 4 to 6 volts; anode current density, 20-60 amp. per sq. ft.

Immersion Tin Solution

Tin Chloride	1/2	oz.
Aluminum Sulphate	2	OZ.
Cream Tartar	2	OZ.
Water	1	gal.

The solution is allowed to boil for 30 to 45 minutes and the addition of a very small quantity of sulphuric acid (about 1 drop to each gal. of solution) hastens the deposition of the tin deposit.

Caustic Soda Method (Tin)

This method is used to tin by immersion, small brass or copper articles.

on copper	CL CLCLCD.
Caustic Soda	12 oz.
Stannous Chloride	4 oz.
Sodium Chloride	1 oz.
Water	1 gal.

The solution is placed in an iron tank, which is heated with a steam coil. The bottom of the tank is covered with moss tin over which is placed an iron wire screen. The work to be tinned is bright dipped or tumbled clean, placed in brass wire baskets and separated with sheets of perforated tin, placed in solution at boiling temperature for 15 to 30 minutes, or until covered with tin. Rinse thoroughly in clean cold water, hot water, dry in sawdust.

Protecting Tin and Lead Against Corrosion

French Patent 777,314

Dip in following solution:	
Copper Sulphate	25 g.
Nickel Sulphate	15 g.
Ammonium Molybdate	3 g. 1 l.
Water	1 l.

Tungsten Plating

The Carbonate Bath:

Tungstic Acid Sodium Carbonate	125 330			
Use at 90° C., 50 amp.	per	sq.	ft.	

The Phosphate Bath:

Tungstic Acid	100 g. per l.	
Sodium Phosphate		
(Na ₂ PO ₄ ·12H ₂ O)	500 g. per l.	

Use at 90° C. with 50 amp. per sq. ft.

Citric Acid Bath:

Tungstic Acid	100	g.	per l.
Potassium Hydroxide	70	g.	per 1.
Citric Acid	250	cc.	per 1.
2.5 Molar Citric Ac	eid)		-

Use platinum anodes; 50 amp. per sq. ft. at 20° C.

Electrolytic Surface Treatment of Zinc

British Patent 421,696

Zinc and alloys consisting mainly thereof are provided with an insoluble coating resistant to weathering and corrosion by anodic treatment in a substantially neutral electrolyte containing an alkali metal ferrocyanide, ferricyanide, dichromate, oxalate or molybdate or ammonium oxalate or molybdate or more than 1 of these. Suitable baths contain 35 g. crystal ammonium oxalate or 50 g. crystal potassium ferrocyanide per l. The metal surface may first be cleaned by cathodic treatment in a bath containing 45 g. sodium phosphate, tribasic, per 1. The coatings may be painted, lacquered or dyed, color coatings being obtainable by adding a dye to the electrolyte.

Zinc Solutions Acid Zinc Solution

Zinc Sulphate	32 oz.
Ammonium Chloride	2 oz.
Sodium Acetate	2 oz.
Water	1 gal.

Temperature 80° F. Cathode current density, 15-20 amp. per sq. ft.; 3-4 volts; pH, 3.5-4.5, using thymol blue as an indicator.

Cyanide Zinc Solution

Zinc Cyanide	4 oz.
Sodium Cyanide	4 oz.
Caustic Soda	3 oz.
Water	1 gal.

Temperature 100° F. Cathode current density 10-15 amp. per sq. ft.; 2-3 volts; keep free cyanide equal to metal content. Use pure zinc anodes. Finish work by rinsing in cold water, then hot water, then drying in hardwood sawdust.

Zinc Cadmium Alloy Plating

Zinc Sulphate	295 g. per l.
Cadmium Sulphate	50 g. per l.
Aluminum Sulphate	30 g. per l.
Caffeine or Licorice	5 mg. per l.

Sulphuric acid may be used in small amounts, but as a general rule, the deposit will not be as bright if acid is present, although appreciably harder. This alloy coating can be deposited directly upon iron, steel, brass, bronze, copper, etc.

Coloring Zinc Dark Brown U. S. Patent 1,853,323

Zine or die cast zine can be colored dark brown by treating in a bath containing:

Chromic Acid 200 g. per l. Sulphuric Acid 2 g. per l. provided the material is treated with an alternating current.

Cleaner for Barrel Plating

Water	1 gal.
Soda Ash	6 oz.
Caustic Soda	2 oz.

This is not suitable for work which has soldered or tinned parts. Such parts should be cleaned in a cleaner which does not readily attack solder or tin. This should be used, 8 oz. to each gal. of water. More may be used without any bad effect upon such work immersed not more than 20 minutes, which will ordinarily clean almost any "hard to clean" parts. It is understood of course that the solution should be kept hot, 180° F.

This cleaner does not readily tarnish brass and copper and has a considerable amount of insoluble material in it which has a scrubbing effect when boiling. This is very effective also in removing oils and dirt and does not require frequent replenishing.

This cleaner is sold on the market under various trade names, the only difference being in the proportions of the 3 sodium compounds.

Another effective cleaning solution used hot or boiling is composed as follows:

Water		1 gal.
Soda Ash		4 07.
Caustic Soda		2 oz.
Trisodium Phosphate		2 07

This too may be varied to suit almost any requirement in cleaning, but a solution made up weaker than the above formula will not work well long. The formula approximates very closely many proprietary cleaners now on the market.

One of the best and simplest combinations for an electrical cleaner is as follows:

Water	1 gal.
Soda Ash	2 oz.
Caustic Soda	1 oz.
Trisodium Phosphate	1 oz.

This may be modified to meet almost any problem of cleaning with the current.

Cleaning Enamel from Metals

Using 50 amp. per sq. ft. at 2½ volts, reversing polarity at 10-second intervals and using following bath gives excellent results.

Caustic Soda

13.6 oz.

 $\begin{array}{cccc} {\rm Trisodium~Phosphate} & 6.38 & {\rm oz.} \\ {\rm Sodium~Silicate} & 1.62 & {\rm oz.} \\ {\rm Water} & {\rm to~make~1} & {\rm gal.} \end{array}$

Cleaning Phosphor Bronze Sheets

After the regular sulphuric acid pickling, they are treated in a bath made of a 10% solution of sulphuric acid with 1½ to ½ lb. of sodium bichromate added to each gal. of the solution.

The general practice is to heat the

solution with live steam.

Metal Cleaner for Electroplating
Sodium Metasilicate 2 lb.
Trisodium Phosphate 2 lb.
Soda Ash 2 lb.
Rosin Soap 0.18 lb.

The quantities given are for each gal. of water in the cleaning tank. Have the water near the boiling point and add the materials by dusting on the surface and stirring until dissolved.

Electroplating Radiators British Patent 425,846

Copper cynanide 40, sodium stannate 20, total sodium cyanide 65, sodium hydroxide, 7.5 g. per l. is specified, this having a free sodium cyanide content of 20 g. per l. and pH 13. A current density of 1–80 amp. per sq. ft., or higher, and a bath temperature 15–17° C. are used. A deposit containing 13–16% tin is obtained. A suitable alloy for automobile radiator shells is tin 15% and copper 85%. The anode preferably consists of an alloy in the proportions of the desired deposit but these may vary by 10% or more. The anodes should be heat-treated to obtain a maximum softness by casting in a metal mold, cooling in the mold,

heating to 1000° F. for 15 minutes and quenching in water. The alkalinity of the bath should be maintained at a pH 12.8-13.5 and the free sodium cyanide at 10-45 g. per l.

Coloring Razor Blades Blue

After blades have been hardened and drawn and being sure that surfaces are absolutely clean, polish well and heat to 550-600° F. This temperature will not affect temper.

Protection of Magnesium by Means of Selenium Coatings

Of many methods tried for coating magnesium with selenium, the following give the same results: (1) immersion for 3 hours in an aqueous solution of 8% sodium selenite, 3.2% selenious acid and 0.10% sodium chloride at 80-90° C; (2) a 10% selenious acid solution with 0.1-0.5% sodium chloride for 5-10 minutes; (3) a 2% sodium selenite solution with 0.2% phosphoric acid for 1 minute; (4) initial cleaning for 30 seconds in 1% chromic acid at 80° and then treatment as in method 3; (5) cleaning as in method 4 followed by method 2.

Increasing Life of Graphite Electrodes

To increase their resistance to attack during electrolysis anodes 25×25 mm. in size are soaked in coal tar for $1\frac{1}{2}-2$ hours at $150-180^{\circ}$ F., or in pitch for 3-5 hours at $300-350^{\circ}$. They are then heated at $300-500^{\circ}$ to drive out the more volatile compounds. Larger anodes require longer treatment. Such anodes are more stable and more efficient than anodes treated with linseed oil. Mixtures of tar and pitch, or bakelite lacquers, may also be used.

POLISHES, ABRASIVES

Aluminum	Pol	ish
Formula	No.	1

Potassium Hydroxide	40 g.
Water	900 cc.
Olive Elaine	150 cc.
Alcohol	25 cc.
Ethylene Dichloride	50 cc.

Add the potassium hydroxide to the water, warm to 75° C. and slowly stir in the olive Elaine until completely dissolved. Cool and add the alcohol and ethylene dichloride.

Directions for Use

Dip a piece of fine steel wool or rough cloth into a liquid and rub on to the aluminum. Then wash the surface with hot water and dry as usual. This aluminum polish used in dish water in proportions of about 2 thsp. per 1 gal. will soften the water and assist in cleaning.

		No. 2		
Whit	ing		75	ğ.
Tripo	li, Fine, Y	ellow	20	g.
	m Bicarbo			3 g.
Potas	sium Sulp	hocyanid	le 2	} ḡ.
Add	Glycerin	Water	(25%)	until
pasty.				

Silver Plating Polish (Renews as it polishes)

Silver Nitrate	30	oz.
Salt	30	oz.
Cream of Tartar	200	oz.
Ont of state of the Land Too	ma oak	***

Grind and sift through 100 mesh sieve. Then make into a paste with

"Celloso Water			parts parts
	Silvar Polish		

Stiver rousii		
Soap	20	oz.
Stearic Acid	5	OZ.
Gilders Whiting	32	oz.
Tripoli	3	oz.
Sodium Thiosulphate	3	OZ.
Water •	37	oz.

Silver Polishing Cloth

<i>a</i> J	Hard Soap Water	10	oz.
۳.)	Water	45	oz.
ъ.	Olein, Distilled	6	oz.

d.	Calcium Carbonate, Precipitated Iron Oxide—Red Ammonia (10%)	5	oz. oz. oz.
		+	oz.
e.	Alcohol	10	٥z.

Dissolve a in an enameled or zincplated or tin-plated steam-heated kettle; when at 60-70° C., add b, stirring to form homogeneous emulsion, then add the powders c. Saponify with d, let stand several hours, and add e. impregnate rags in this solution.

Non-Scouring Copper Polish Make a paste of finely powdered glass and mineral oil. This will not scratch.

Polish for Chromium, Liquid a. Hard Soap, Powder 3 g. b. Water, Hotc. Olein, Distilled 53 cc. 5 cc. d. Ammonia (10%) 3 cc. e. Alcohol, Denatured 16 cc. f. Tripoli 20 g.

Dissolve a and b, saponify with c, dilute with e, add f.

Chromium Polishes

Chromium Ponsner	3	
Formula No. 1		
Olein Stearin	20 60	
Melt. Calcium Carbonate (Powdered) Cool, powder.	20-30	g.
No. 2		
Chromium Oxide . Stearic Acid or	60	g.
Paraffin Wax	40	g.
No. 3		
Carnauba Wax Yellow Wax Japan Wax Paraffin Wax (46-48° C.)	10 15 15 60	g.
Melt on water bath.		
Melt together and add:		
Turpentine Tripoli, Dry	130 70	
Turpentine	100	

until pasty

No. 4	
Rouge (Iron Oxide)	50 g.
Kieselguhr, White, Burned	100 g.
Neuburger Chalk	150 g.
Coconut Oil Soap	700 g.
No. 5	
Chromium Oxide, Powdere	d 50 g.
Paraffin Wax	50 g.
Emery	30-50 g.
No. 6	0
Stearin	90 g.
Stearin Oil	25–30 cc.
Neuburger Chalk	30-45 g.
Melt together.	
Cool; powder.	

Polish for Metals French Patent 772,648

Formula No. 1

A polishing compound contains kaolin 30-50, tale 10-20, rosin 18-30, alcohol 4-15, ammonia 5-18 and acetone 1-10 parts by weight.

No. 2

French Patent 772,691

A compound contains powdered silicon dioxide 35, soap powder 5.9, neutral oil 0.23, ammonium sulphate 3.1 and bentonite 0.63 kilograms.

No. 3			
Kieselguhr	2	pa	rts
Strong Ammonia Water		. pa	
Denatured Alcohol		. pa	
Shake well with water	q.s.	to	give
reemy consistency	•		•

creamy consistency. Metal Polish (Sidol Type)

a. Olein, Distilled 4.5 cc. Stearin ٦ Alcohol čc.

Heat to 50° C.

b. Ammonia (sp. gr. 0.91) cc. Saponify.

c. Oxalic Acid g. Water (50-60° C.) 70 cc. d. Neuburger Chalk g. Optional:—add more water.

Metal Polish Block

	Stearin Olein	25	g.
u.	Olein	5	cc.
7.	Spindle Oil, Refined Vienna Lime	2-10	cc.
0.	Vienna Lime	30	g.
c.	English Red (Ferrous		
	Oxide)	38-30	g.

Mix first b, to prevent saponification of the fats a.

Black Polish for Ovens

Formula No. 1

Graphite, Flaky	1000 lb.
Lampblack	50 lb.
Beeswax, Crude	10 lb.
Montan Wax, Crude	100 lb.
Paraffin Scales (50-52° C.)	30 lb.
Melt together.	
Nigrosine, Fat Soluble	5 lb.

Naphtha

No. 2	
Graphite, Colloidal	20 lb.
Paraffin Wax	13 lb.
Lacquer Benzoline (White	
Spirits)	67 lb.

Ma 9

	740. 9		
a.	Olein, Distillate	15	cc.
	Stearin (52-54° C. titer) 5	g.
	Ammonia (25%)	4	cc.
c.	Spindle Oil	10	cc.
	Alcohol	40 - 50	cc.

Melt a on water bath, saponify with b, add c, then an abrasive (Emery, Carborundum, Chromium Green, Graphite).

Automobile Polish Cleaner

Formula No. 1

	Olein	10	cc.
	Mineral Oil	20	cc.
a.	{ Petroleum	20	cc.
	Turpentine Oil or		
	White Spirit	28	cc.
7.	Alcohol Ammonia (0.910)	6	cc.
о.	Ammonia (0.910)	6	cc.
c.		10	g.

No. 2		
Yellow Wax	10	oz.
*Air-Floated Tripoli	18	oz.
White Spirit	19	oz.
Soft Soap	1/2	oz.
Water	216	OZ.

Melt the wax in a double pan and add the powder slowly; keep stirring while slowly adding the white spirit. Dissolve the soft soap in the water and add to the mix with constant stirring. On cooling this forms a soft paste.

A liquid polish can be made as follows:

No 3

7/0. 9		
White Spirit	$2\frac{1}{2}$	pt.
Mineral Oil	21/2	
Turkey Red Oil	4	pt.
Ammonia	1	oz.
Water	5	pt.
Glycerin	1	pt.
Formaldehyde	. 8	07.

*Fuller's Earth 8 oz. *Bentonite 6 oz.

Mix the oils together first and add the abrasive powders, then the water, ammonia, glycerin, and formaldehyde; stir rapidly until a smooth mixture is obtained.

*The quantity and type of abrasive used can be varied according to whether the polish is to have a strong or mild abrasive action. Polishes to be used as maintenance polishes by car owners should be only mildly abrasive, otherwise too much of the finish will be rubbed off.

Car Polishes Formula No. 1

a. Spindle Oil, Refined 80-85 g. Methyl Hexalin 20-15 g. b. Distilled, Warmed

Water 400-900 g.

Add b to the mixture a with high speed stirring.

Apply spraying and polish with a rag. No. 2

Linseed Oil Dipentene Paraffin Oil Patroloum Pafined	200 g. 300 g. 200 g.
Petroleum, Refined Camphor Oil, Light	250 g. 50 g.
Apply simply with rag.	

Automobile Cleaner and Polish

Kieselguhr	30	oz.
Tripoli	5	oz.
Paraffin Wax	4.5	0 z .
Carnauba Wax	0.5	oz.
Varnolene	30	oz.
Tint with iron oxide		

Automobile Paste Polish

Carnauba Wax	5 oz.
Beeswax	5 oz.
Ceresin Wax	5 oz.
Stearic Acid	2 oz.
Soap	2 oz.
Varnolene	45 oz.
Water	10 oz.

Automobile Polish, Powdered

Mineral Oil	5 lb.
Kerosene	10 lb.
Diglycol Laurate	1 lb.
Silica Dust	½ lb.
Kieselguhr	4 lb.
Tripoli	1 lb.

Automobile Polish (Tumbler's) U. S. Patent 1,969,387

To 31/2 gal. of pale blown castor oil. add 34 gal. of orthodichlorbenzol. This is mixture No. 1. To 15 gal. of water, add 11 gal. of neutral pale mineral oil and ¾ gal. of ammonia, which has been previously made up of one part of ammonia of 26° Bé. and 4 parts of water. This is mixture No. 2. Mixture No. 1 and mixture No. 2 are combined and agitated for about 5 minutes. and one-half gallons of special petroleum spirit is added and the whole mass is now stirred about 10 minutes. It is then run through a colloid mill and is ready for use. Alternatively, all of the ingredients may be mixed in a single batch and passed through the colloid mill, which breaks up the particles to a fine degree. This obviates preparing separate mixtures.

Auto Polish U. S. Patent 1,979,787

Wax Base	
Carnauba Wax Petrolatum Wax (160 to	66.5 g.
165° F. Melting Point) Petrolatum (140° F. Melt-	26.6 g.
ing Point) Rosin	6.3 g. 0.6 g.
Wax Base (Prepared as	0.0 g.
Above Described) Refined Mineral Oil (Nar-	9 g.
row Cut) Starch	41 g.
Water	0.5 g. 49.5 g.

The refined oil is a distillate having an initial boiling point of about 350 and an end point of about 475° F. Although it is not necessary that these precise limits be maintained, it is important that a narrow cut be used of about this range. The so-called "W.W. 150" (water white kerosene), with a boiling range of about 373 to 504° F. evaporates too slowly, while oleum spirits, with a boiling range of about 300 to 425° F. evaporates too rapidly to give best results. The narrow boiling range of the refined oil is of particular importance in a "set" or solid emulsion of this type. It is also of particular importance that the oil be highly refined (treated with sulphuric acid for the removal of unsaturateds and other impurities) because untreated light petroleum distillates may be injurious to the skin.

In preparing the finished product melt

the base stock with the refined oil and heat the mixture to a temperature of about 175 to 200° F. Then boil a 1% starch solution and make an oil-in-water emulsion in a colloid mill at a temperature above the melting point of the water, usually at about 130 to 200° F. When the resulting emulsion cools, it sets to form a semi-hard, solidified emulsion which is extremely stable and which possesses entirely different structural properties from the ordinary liquid oil-inwater emulsions of the same concentrations. The product may be stored for an indefinite period of time without separation, and it may be easily handled and applied.

Solid Abrasive Polish (Automobile	Wax),
Formula No. 1	•
a. Montan Wax, Bleached Paraffin (40-42° C.) Ozokerite, Refined	8 g. 8 g. 2 g. 35 g. 13 cc.
b. { Infusorial Earth Spindle Oil, Refined White Spirit c. Turpentine Oil or Substitute	35 g. 13 cc. 13 cc. 21 cc.
No. 2	
a. Montan Wax, Bleached Montan Wax, Double Bleached Olein	8 g.
a. Bleached Olein	5 g. 2 cc.
b. Potassium Carbonate Glycerin (28° Bé.) Water, Boiling	2 g. 3 cc. 40 cc.
c. Yellow Clay or Bentonite	to suit
d. Turpentine Oil or White Spirit	22 cc.
Melt a, add hot (boiling) cool, add d.	b, then c;
* ·	
Auto Polish	
Formula No. 1	-

Auto Polish		
Formula No. 1		
Montan Wax, Bleached	4	g.
Paraffin Wax (50-52° C.)		g.
Hard Soap		g.
Water		cc.
Water Soluble Dyestuff	2	g.
(Black: 4 parts Nigrosine)		
Ammonium Hydroxide (0.910		
Alcohol No. 2	20	cc.
Montan Wax, Bleached		1.
Soft Soap	3	g.
Potassium Carbonate	0.8	g. g.

DRASIVES	28
Water Water Soluble Dyestuff (Black: 4 parts Nigrosine	87.2 cc. 2 g.
No. 3	
Shellac (Orange) Alcohol Carnauba Wax Paraffin Wax (50-52° C. Turpentine	14 g. 60 cc. 2 g. 1 g. 23 g.
Polish for Lacquered or Poli Swiss Patent 172,7	shed Object
Turpentine Paraffin Beeswax	100 cc. 50 g. 15 g.

Turpentine	100 cc.
Paraffin	
Beeswax]
Silica Powder	δ.
Chalk Meal	_ 5.
Vienna Lime	2 0.
Oxalic Acid	
Ammonia (28%)	1 g.
(2070)	10 cc.

Polish for Leather Furn	iture
Paraffin Wax (50-52° C)	20 ~
Ozokerite/Ceresin (58-60° (Wool Fat, Neutral	
Beeswax	5 g. 10 g.
Carnauba Wax Turpentine Oil	10 g.
Color similar to that of	150 g.
Pour at 40-45° C. into jar.	- armiture.

Furniture Polish Formula No. 1 Raw Linseed Oil 10 oz. Spindle Oil 50 oz. Stoddard Solvent 15 oz.

04-33- 3.0 3			oz.	
Stoddard Sol	vent	15	oz.	
Xylol			oz.	
Soft Soap			oz.	
Water				
11 6001		19	OZ.	
	No. 2			
Paraffin Oil		90	0.77	

Paramin Oil	20 oz.
Red Oil	5 oz.
Soft Soap	
	3 oz.
Gum Arabic	2 oz.
Water	70 oz.
emi .	10 02.

The above are mixed vigorously until completely emulsified.

No. 3	
Carnauba Wax	2 g.
Montan Wax, Bleached	6 g.
Beeswax	5 g.
Paraffin Wax (52-54° C.)	14 g.
Melt.	5.
Add:	
Linseed Oil (or Varnish)	3 g.
And (when temperature is	43-45° C.)

Turpentine 70 g.

	1
Liquid Furniture Polish	At same time prepare:
	c. Potassium Carbonate 5 g.
a. Beeswax, Yellow 13 g. Ozokerite, Yellow 2 g.	Hard Soap 5 g.
b. Thinner (White Spirit) 75 cc.	Water, Hot 45 cc.
c. Alkali Solution (Water: Am-	and pour in thin jet into a plus b, stir.
monia (0.91) = 85:15) 10 cc.	Keep temperature at 55-60° C. Stir con-
Melt up a , add the warmed b to clear	tinuously, add a yellow dye, then pour
solution, then add c in thin jet, stirring	into cans.
thoroughly.	No. 2
unoroughij.	Paraffin Scale 12 g.
Furniture Polish	Shellac Wax 5 g.
Formula No. 1	Carnauba Wax 4 g.
a. Paraffin Oil, Yellow 100 cc.	Ozokerite Ceresin (58-60° C.) 3 g.
Naphtha, Refined 50 cc.	Montan Wax, Bleached 4 g.
Tetralin, Dipentene 50 cc.	Turpentine Oil Substitute 72 cc.
Precipitated Chalk 25 g.	No. 3 (White)
b. Lactic Acid (50%) 50 cc.	1
Water 225 cc.	Carnauba Wax, Bleached 6 g. Ozokerite, Refined 4 g.
Add b to a in thin, continuous jet;	Ozokerite, Refined 4 g. Paraffin (50–52° C.) 20 g.
stir well.	
No. 2	Thinner (Turpentine Oils, Dipentene, Hydroterpene, Dec. 70 g.
Boiled Linseed Oil 10 lb.	aline, White Spirit)
Raw Linseed Oil 12 lb.	
Denatured Alcohol 2 lb.	No. 4 (White)
Vinegar 12 lb.	Montan Wax, Double
Turpentine 14 lb.	Bleached 12 g.
Petroleum Spirits 27 lb.	Montan Wax, Bleached 5 g.
or	Paraffin $(50-52^{\circ} C_{\bullet})$ 6 g.
Raw Linseed Oil 2 gal.	Ozokerite, Refined 2 g.
Paint Drier ½ gal.	Thinner 75 g.
Vinegar 6 gal.	No. 5 (White)
Furniture Finishers Polish	Montan Wax, Double Bleached 8 g.
Turpentine 7 lb.	Bleached 8 g. Montan Wax, Bleached 3 g.
Mineral Oil 7 lb.	Paraffin (50–52° C.) 19 g.
Cedarwood Oil 2 oz.	Thinner 70 g.
Sassafras Oil 1 oz.	
Rottenstone, Fine Powdered 4 oz.	No. 6 (Yellow or Orange)
Programme of the Control of the Cont	Carnauba Wax, Fat-Gray* 4 g.
Covering Polish for High-Gloss Polished	Ozokerite, Yellow 2 g.
Furniture	Paraffin (48-50° C.), Yellow 24 g. Thinner 70 g.
Collodion Wool (Nitrocellu-	
lose), Alcohol Soluble, 12 g.	* Dye with 0.02% Sudan Yellow G.
soaked in Butanol (2:1)	• 77.77
Ethylene Glycol 6 g.	Liquid Floor Polish
Toluene 12 g.	Melt:
Tricresyl Phosphate 2 g.	Paraffin Wax (50-52° C.) 50 g.
Shellac (Free from Wax) 10 g.	Ceresin (58–60° C.) 10 g.
Alcohol (95-96%) or	Carnauba Wax 40 g.
Butanol 58 g.	and dissolve:
Thinner (Alcohol) optional	In summer, 7-9 parts in 93-91 parts
X X X X X X X X X X X X X X X X X X X	of turpentine.
Floor Polish	In winter, 6-7 parts in 94-93 parts of
Formula No. 1	turpentine.
Carnauba Wax 15 g.	
a. Montan Wax 5 g.	Deodorized Floor Polish
Rosin, Pale 5 g.	TO . 00 TIT. (50 500 C) TO
Melt on water bath, put out fire. Add:	7 7 777
b. Turpentine Oil, or	0 (50 000 0)
Substitute 20 cc.	Rosin, Pale 4 g.
	. 5.

Stearin Potassium Carbonate Caustic Soda (38° Bé.)	1 g. 2 g. 0.5 cc.	No. 2 Spindle Oil, Refined (see
	6.5 cc.	above) 60 cc.
Boil and stir until smooth.		Petroleum 27 cc. Camphor Oil 3 cc.
Managed and the state of the st		No. 3
Dyestuffs for Floor Polis	shes	Spindle Oil, Refined (see
Yellow:		above) 50 cc.
Sudan Yellow RRN		Benzine 40 cc. Turpentine 5 cc.
Orange: Sudan Orange G, RR		Citronella Oil 5 cc.
Red: Sudan Red 5B		Mop Oil Polishes
Brown:		Above given formulae, but adding
Sudan Brown B, 3B, RRN		Waxes (as Montan Wax, Bleached, or Paraffin Scale
Reddish Brown: Sudan Brown 3B	66%	Wax) 2-3 g.
Sudan Red 5B	34%	Dye with
Chocolate Brown:	70	Sudan Dyes 0.02 g.
Sudan Brown 3B	60%	or
Sudan Red 5B	30%	Basic Dyes 0.06 g.
Sudan Black BT	10%	Weden ((C.1.11.11.77)
Other Oil Coloring Base	S	Water "Soluble" Floor Oil
Yellow: Leather Yellow—Fat Dye		Spindle Oil, 5E (20° C.) 40 cc. Tallöl, Crude 20 cc.
Orange: Leather Yellow—Fat Dye	66%	Mix, warm to 70° C., add in thin jet:
Red Fat Dye	34%	Caustic Soda, 38° Bé. 8 cc.
Red:	,,,	Boil to saponify, add again
Red Fat Dye		Spindle Oil (as above) 27 cc.
Brown:		Boil shortly, add boiling Water (to thin the alkali) 5 cc.
Brown Fat Dye		Use: 1 part oil in 6-10 parts water.
Reddish Brown: Brown Fat Dye	66%	- Doc. 1 part of in 0 10 parts water.
Red Fat Dye	34%	Yellow Floor Wax
Chocolate Brown:	/*	Formula No. 1 No. 2 No. 3
Brown Fat Dye	60%	Paraffin Wax 16000 16000 16000 g.
Red Fat Dye	30%	Carnauba Wax 3000 3000 2500 g.
Ceres Black I, pieces Pigments:	10%	Beeswax, Yellow 1000 2000 1500 g.
Red: Iron Oxide Red		Turpentine 46000 25000 30000 cc.
Brown: Iron Oxide Brown		(Dye) 20 20 20 g.
		Amyl Acetate — — 100 cc.
Floor Oils		Million and the state of the st
Spindle Oil, Pale, Viscosity		Dance Floor Wax
2.5-5E (20° C.), Ignition Point 160-200° C.	95 сс.	Formula No. 1
Olein	5 cc.	Melt
	0 00.	Paraffin Scale (Yellow, 50–52° C.) 12 g.
Mop (Floor) Oils		a. Yellow, 50-52° C.) 12 g. Dye, Yellow or Red,
Formula No. 1		l Oil Soluble 25-30 g.
Spindle Oil, Refined.		b. Talc 80 g.
sp. g. $= 0.850$; 1.8–2.5E (20° C.)		Ochre, Yellow 8 g.
(20° C.)	70 cc.	Mix a and b thoroughly, cool, pulverize.
Benzine Balm-Turpentine Oil, Hydro-	25 cc.	Melt No. 2
terpene, Wood-Turpentine		Paraffin Wax (50-52° C.) 80 g.
Oil or Refined Pine Oil	5 cc.	Carnauba Wax, Refined 20 g.

Sudan Yellow Sudan Red	}	to	suit
3.5-14. 4 43	T		

Melt together, cool, pulverize.

Linoleum Wax

The following waxes are suitable for preservation of linoleum. The clear wax is also suitable as a floor wax or as a polish.

Clear Wax

Carnauba Wax	1 lb. 6 oz.
Ceresin Wax	1 lb. 6 oz.
Petroleum Spirits	8 lb.

Melt the two waxes together and stir in the petroleum spirits. The wax should then be ground.

Red Wax

Carnauba Wax	11/2 lb.
Ceresin Wax	1½ lb.
Venetian Red, Dry	1/2 lb.
Petroleum Spirits	61/2 lb.

Red Stain for Linoleum

Venetian Red, in Oil	$1\frac{1}{2}$	lb.
Boiled Linseed Oil	3 -	pt.
Amyl Acetate	41/2	pt.

Wax Polishes

U. S. Patent 2,010,297

rormu	na no. 1	140.	~
Carnauba Wax	25 g.	2.75	g.
Ceresin Wax	28 g.	3.08	g.
Beeswax, Yellow	20 g.	2.20	g.
Montan Wax	22.5 g.	2.47	g.
Calcium Stearate	4.5 g.	0.50	g.
Light Petroleum	•		_
Solvent	<u> </u>	89	g.

The four waxes should be melted together at about 200° F., or somewhat higher, and the calcium stearate then dissolved in the molten wax with gentle agitation. When the melt becomes clear, about half of the solvent is added. The solution is then cooled, to as low a temperature as 135–140° F. and vigorously agitated as by means of high speed stirrers, with the cooling continued until crystallization occurs around 100–110° F. The vigorous agitation is further continued until the batch reaches a temperature of 90–95° F., whereupon the other half of the solvent is slowly added in connection with gentle agitation. The product may then be packaged.

Wax Paste Polish

Paraffin				28	ø.
Ozokerite					g.
Carnauba	Wax, N.C	No.	3		g.
Beeswax,	Yellow				g.
Turpentin	ne			60	čc.

I	Emery	Polishing	Paste	

Emery, Powdered	45 g.
Aluminum, Powdered	4 g.
Wax Paste Polish	24 g.

Wax Polish U. S. Patent 1,979,787

Carnauba Wax	9	lb.
Light Petroleum Oil	41	lb.
Water	49.5	lb.
Starch	0.5	lb.

Wood Button Polish

Turpentine	120 cc.
Wax, White	120 g.
Melt.	
Add Alcohol with stirring.	50 cc.
with stilling.	

Axe or Hammer Handle Wax

White Beeswax	51/2	lb.
White Rosin	1/2	lb.
White Lead	4	lb.
Damar Varnish	1/s	lb.

Melt the beeswax; crush, melt and stir in the rosin; add white lead while stirring, and finally pour in the damar varnish. While still in a liquid state, this material is poured into small paper bags which serve as molds.

Another mixture contains:

10	lb.
2	lb.
2	OZ.
1/2	lb.
Controlled.	10 2 2

The finished product looks like beeswax, but is lighter in color. The rosin and paraffin are melted and mixed and allowed to cool somewhat before stirring in the white lead and linseed oil—this to prevent foaming.

Liquid Ski "Waxes"

	rormura	77.00	1		
Shellac				90	2.
Sandarac				10	
Alcohol				200	

Use solution to spread over the lower surface of the ski, from the top down, to about 10 cm. below the straps. Dry, and repeat spreading. For low temperatures, when snow has too much friction, add 1-2% Castor Oil.

No. 2

Carnauba Wax		4	g.
Montan Wax		12	
Linseed Oil Varnish		84	

No. 3		No. 6	
Montan Wax, Refined	15 g.	Montan Wax, Crude	120 g.
Ceresin	3 g. 82 g.	Paraffin	30 g.
Turpentine Oil Substitute	02 g.	Wool Fat Seal Train Oil	20 g. 15 g.
No. 4	20	Tallow, Hard	10 g.
Colophony Ceresin	30 g. 25 g.	Rosin	5 g.
Tallow	55 g.	Wood Tar	3 g.
No. 5		No. 7	1 g.
Tale Palm Oil	16 g. 14 g.	Tallow	1.5 g.
Ceresin	16 g.	Rosin	$_{2.5}^{2.5}$ g.
Paraffin	60 g.	Ozokerite No. 8	15 g.
No. 6	125 g.	Wool Fat	10 g.
Tallow Colophony	275 g.	Ceresin	90 g.
Montan Wax	400 g.		•
Turpentine Oil	200 g.	Ski Wax	
No. 7		Formula No.	1
Rice Starch	40 g. 125 g.	Montan Wax, Crude	18 g.
Tallow Larch Turpentine	260 g.	Paraffin Wax	60 g.
Yellow Wax	500 g.	Ozokerite	4 g.
No. 8		Wool Fat Colophony	6 g. 12 g.
(Sohm's Ski Wax)	Melt together and add	
Ozokerite	55 g.	to desired consistency.	tarpontine on
Tallow	15 g.	27. 0	
Rosin	30 g.	No. 2	
All these waxes may be to turpentine oil to desired fluid		Ascension Wax:	10 m
tarpointino on to decirate	, -	Paraffin Wax	10 g. 20 g.
		Wool Fat	28 g.
Norwegian Klister (Ski)	Waxes	Colophony Montan Wax	15 g.
Formula No. 1		Melt together and add	27 g.
Rice Starch	40 g.	desired consistency.	rarbearine co
Tallow	125 g. 260 g.	*	
Larch Turpentine Yellow Wax	500 g.	No. 3	
No. 2	J	Gliding Wax: Paraffin Wax	co
Paraffin (40-42° C.)	60 g.	Ceresin Wax	60 g. 16 g.
Colophony	12 g.	Tallow	14 g.
Wool Fat Carnauba Wax	6 g. 4 g.	Melt together and add	turpentine to
Montan Wax	80 g.	suit.	•
No. 3		Gliding Wax:	
Ozokerite	55 g.	Black Ozokerite	55 g.
Colophony Spindle Oil, Refined	35 g. 10 g.	Rosin	30 g.
A STATE OF THE STA	10 8.	Tallow	15 g.
No. 4	70 m	Melt together and add	turpentine to
Paraffin Colophony	70 g. 15 g.	Suit.	
Wool Fat	10 g.	Paraffin Wax	30 g.
Carnauba Wax	5 g.	Montan Wax, Bleached	80 g.
Montan Wax No. 5	15 g.	Colophony	20 g.
Ozokerite	5 g.	Japan Wax Wood Tar Oil	20 g. 10 cc.
Colophony	4 g.	Turpentine Oil	10 cc.
Train or Spindle Oil	1.5 g.	Yellow Dyestuff eno	ugh to color

No. 6		High-Luster Polish for 8	Shoes
Wax Polish, White:		Formula No. 1	
Paraffin Wax Carnauba Wax, Light Beeswax, White Turpentine No. 7	16 g. 3 g. 1 g. 46 cc.	Carnauba Wax, Yellow Carnauba Wax Residue a. Montan Wax, Bleached Paraffin (50-52° C.)	500 g. 500 g. 500 g. 200 g. 150 g.
Wax Polish, Liquid: Paraffin Wax Ozokerite Carnauba Wax Turpentine Oil Benzoline Camphor Oil Amyl Acetate	50 g. 5 g. 100 g. 750 cc. 94 cc. 2 g. 3 cc.	b. Potash, Caustic Olive Oil Soap c. Turpentine Oil or	8500 cc. 300 g. 100 g. 1500 cc. , stir until
No. 8		No. 2	
For Gliding: Paraffin (50-52° C.) Ceresin (60° C.) Tallow or Palm Oil Talcum No. 9	60 g. 16 g. 14 g. 10 g.	Montan Wax, Crude Carnauba Wax Ozokerite (58–60° C.) Candelilla or Shellac Wax Paraffin Scales (50–52° C.)	6 g. 3 g. 2 g. 3 g. 14 g.
For Climbing:		Nigrosine Base Turpentine 2	3 g. 30–30 cc.
Paraffin (40–42° C.) Rosin Wool Fat Wood Tar No. 10	50 g. 20 g. 15 g. 15 g.	Shoe Polish Paste Carnauba Wax, Fat-Gray	6 g.
Climbing and Sliding Ski W Paraffin Montan Wax, Crude Wool Fat, Neutral Rosin Mineral Oil Wood Tar	40 g. 15 g. 15 g. 10 g. 15 g. 5 g.	Montan Wax, Bleached Paraffin (50-52° C.) Ozokerite Dyestuff Thinner (Turpentine Oil, or substitute or a Mixture of Both)	7 g. 11 g. 2 g. 2 g.
No. 11 Climbing Wax:			
Montan Wax, Crude Wool Fat, Neutral Paraffin Rosin Ozokerite Mineral Oil Wood Tar	17 g. 18 g. 10 g. 28 g. 25 g. 5 g. 2 g.	Shoe Polish British Patent 395,55 Paraffin Ozokerite Carnauba Wax Melt 80-90° C. Turpentine Oil	38 14 g. 3 g. 3 g. 38 cc.
		Stir now with	
Ski Finishes For running on wet snow.		Water (boiling) Sodium-Sulphonate of Glycol Mono-Oleate	
Mix: Pine Tar Copal Lacquer Venice Turpentine	25 g. 25 g. 50 g.	Dyeing Shoe Polish, Lie	1 g. quid
This mixture is boiled in ning side of the ski with a Before using the ski rub in ing of Venice turpentine. For running on very cold in a good coating of Pine tarusing heat ski and rub on aceti.	on the run- a blowtorch, a thin coat- snow burn r and before	Carnauba Wax, Fat-Gray Montan Wax, Bleached Paraffin (50-52° C.) Ozokerite, Refined Dyestuff Thinner (Turpentine Oil, or Substitute, or Mixture of Both)	2 g. 2 g. 4 g. 1 g. 1.5 g.

Sporting Shoe Dressings, Formula No. 1 Shoe Paste, Black Carnauba Wax, Gray Montan Wax, Crude Paraffin (50-52° C.) Black Dye, Oil-Soluble * Thinner (Turpentine Oil, or Substitute, or a Mixture or Both) Vaseline Oil No. 2 Carnauba Wax Montan Wax, Crude	7 g. 7 g. 12 g. 3 g. 8 g. 20 cc. 5 g. 8 g.	Paraffin (50-52° C.) Black Dye, Oil-Soluble * Thinner (see above) Spindle Oil, Refined No. 3 Carnauba or Shellac Wax Montan Wax, Crude Paraffin (50-52° C.) Black Dye, Oil-Soluble * Thinner (see above) Spindle Oil, Refined Sardine Train Oil	9 g. 3 g. 60 cc. 15 cc. 8 g. 8 g. 10 g. 3 g. 51 cc. 10 cc.
Spo	rting Shoe	Polishes, Liquid	

Sporting Shoe Polishes, Liquid	Sporting	Shoe	Polishes.	Liquid
--------------------------------	----------	------	-----------	--------

	Formula.	No. 1	No	. 2	No. 3
Carnauba Wax, N.C.		3 g.	3	g.	4 g.
Montan Wax, Crude		2 g.	2.5	g.	2 g.
Paraffin (50-52° C.)		3 g.	2.5	g.	2 g.
Black Dye,* Oil-Soluble		3 g.	3	g.	3 g.
Thinners (see above)		75 cc.	72	cc.	70 cc.
Spindle Oil, Refined		14 cc.			19 cc.
Vaseline Oil			11	cc.	
Sardine Train Oil			6	cc.	

* Black Dyes	
Nigrosine Base	51017
Nigrosine Base	4322
Nigrosine Base	LJF
Nigrosine Base	srn
Nigrosine Base	sr
Nigrosine Base	C
How to dissolve the Black Dye	:
a. Olein	1 g.
Montan Wax, Raw	1 g.
Nigrosine Base	1 g.
Warm together and stir.	
or	
b. Stearin	2 g.
Nigrogina Raga	1 or

Nigrosine Base	1 g.
Black Shoe Polisi	h
Commonly War	6 ~

Carnauba wax	og.
Montan Wax, Crude	5 g.
Soft Ozokerite (58-60° C.)	1 g.
Nigrosine Base	3 g.
Paraffin (58-60° C.)	14 g.
Turpentine	71 cc.
*	

Powder Glaze for Shoes

Shellac			18	g.
Borax			$7\frac{1}{2}$	g.
Water			75	g.
Diagolmo	d ther	0770700		TTTO to

Dissolve and then evaporate until dry and then pulverize.

Shoe Cream for Collapsible Tubes

a.	Water	52	cc.
	Nigrosine	1	g.
	Potassium Carbonate	e 0.5	g.
	Hard Soap	0.75	g.
	Boil.		

	0 00.		
ъ.	Montan Wax, Crude	7	g.
	Japan Wax	2.5	g.
	Carnauba Wax, Gray	4	g.
	Beeswax	2.5	g.
	Paraffin (50-52° C.)	2	g.
	Oil-Soluble Black	2.5	ø.

Pour b molten into hot a. homogeneous (cooled) mass add while stirring

c. Turpentine

Notes on Cleaning White Shoes

Important note—all cleaners should be applied sparingly. It is best to place the shoes to be cleaned on the shoe trees and with a dry cloth remove surface dust or dirt. Do not clean white shoes while on the feet.

Apply the cleaner sparingly to a clean white cloth, preferably toweling, and first clean the dirtiest spot, then go all over the shoe, using sufficient pressure to remove all spots and stains. Avoid saturating the leather but apply evenly over the entire area to be cleaned.

Permit shoes to dry thoroughly. Next rub the shoe briskly with a clean dry cloth, removing all white particles of powder and until the original sheen is restored.

In the case of white buck or suede shoes, a fine bristle brush will more easily remove excess powder and raise the nap of the leather.

Do not use soap and water on elk shoes. Beware of a cleaner with so much alkali that repeated usage will remove the finish. This generally results in the hardening of the elk leather so that it cracks or shrinks.

White Shoe Polishing Stick

Carnauba Wax, Flora		4 lb.
Stearic Acid		4 lb.
Paraffin Wax	,	17 lb.
Montan Wax, Bleached		16 lb.
China Clay		9 lb.
Titanium Dioxide		1 lb.

White Shoe Dressing

Titanox A	10.5	oz.
Titanox B	20.75	oz.
White Soap	3	oz.
White Dextrin	3	$0Z_{\bullet}$
Ammonia	1.25	oz.
Water	48.40	oz.
Carbon Tetrachloride	13.25	oz.
Moldex or Other		
Preservative	10	oz.

Shoe White (Water Type)

This cleaner for white canvas and leather shoes cleans and whitens at the same time and leaves a coating which does not dust or rub off.

Li	ithopone	28	oz.
A	sbestine	4	oz.
G	ım Arabic		oz.
	um Tragacanth		oz.
\mathbf{B}	enzoate of Soda or Mold		
	ltramarine sufficien	t to wh	iten
P	erfume		

sufficient to give pleasant odor Water 59.7 oz.

If better hiding power is desired titanium dioxide pure or titanium dioxide with a barium or calcium base may be used; as well as pure zinc sulphide. The asbestine is added to prevent the pigment from packing hard on long standing. The tragacanth gives added body or viscosity, and inhibits much of the pigment from settling, a mere inversion of the bottle being adequate to bring same back into suspension.

Shoe White (Waterproof Type)

This composition leaves a coating which is waterproof and does not dust off. It is preferred to the water type for leather shoes particularly the glazed type.

Lithopone	28 oz.
Asbestine	4 oz.
Ultramarine sufficie	ent to whiten
Ester Gum, Pale	5 oz.

Solvent Naphtha Aluminum Stearate	-	oz.
Perfume		

sufficient to mask petroleum odor

The solvent naphtha should be a petroleum fraction boiling between 200° and 300° F. The aluminum stearate is dissolved in same to increase the viscosity and inhibit settling of the pigments. The ester gum is then added and stirred or heated until solution is complete. The perfume and pigments are then added.

White Shoe Cleaner

Titanox C Diglycol Laurate Varnolene	30 6 10	cc.
Toluol	12	

Mix a and b thoroughly.

	Drying Carnauba		
Wax	Emulsion	60	cc.
Water		20	cc.

Add c to ab in 4 equal portions, shaking or stirring during and after each addition.

d.	Trichloroethylene	40	cc.
Ad	d slowly with stirring.		

White Shoe Dressing

Titanium White	6	0 g.	
Diglycol Oleate		2 g.	
Naphtha	2	0 g.	

Stir the above together and while stirring vigorously add slowly

Carnauba Wax Emulsion

(10% Wax)	80	g.
then stir in vigorously		
Trichloroethylene	60-100	g.

Polishing Cloths

Prepare powder mixtures: Formula No. 1

- 0122202	
Calcium Carbonate	70 g.
Kieselguhr	25 g.
Caput Mortuum	5 g.

No. 2

Magnesia, Calcined English Red Vienna Lime		$\frac{20}{40}$	g.	
No. 3				

Calcium Carbonate 40 g. Bolus 20 g. Vienna Lime 20 g.

Infusorial Earth		10 g.
Magnesia Usta		5 g.
One hundred and	fifty gram	s of these

mixtures are stirred into 1000 cc. of

water, impregnate the cloths in this suspension. Press. Dry (40-50° C.). Fix with a bath of 100 g. hard soap in 1000 cc. water. Press and dry again.

Cleansing and Polishing Compositions British Patent 425,323

A cleansing and polishing liquid which leaves a thin film on the leather, wood, metal, or other article treated, is composed of a hard wax polishing composition, alkali, water, a solvent of oil and fat, carbon tetrachloride shellac, and bornyl acetate. For example, 3 lb. of shellac wax, 3 lb. of montan wax, 3 lb. of carnauba wax, 2 lb. of paraffin wax, 1 lb. of japan wax, 1 lb. of acetone varnish, 1 lb. of nitrocellulose varnish, 1 lb. of cellulose varnish, 3 lb. of potash or soda, 20 lb. of water, 1 lb. of castor oil, 5 lb. of white spirit, 40 lb. of turpentine substitute, 20 lb. of carbon tetrachloride, 1 lb. of shellac, and 1 lb. of bornyl acetate are mixed together.

Pore Filler for Polish Bases German Patent 607,521

Carnauba Wax	5	g.
Pumice Powder	100	ğ.
Sandarac	100	ġ.
Castor Oil, Blown	10	g.
Shellac Wax	10	g.

Melt up while stirring, cool, and pulverize. The "Pore Filler" is then ap-

plied as usual by rubbing it in on the wood surface together with the polishing liquid.

Abrasive Wheel Formula No. 1 British Patent 411,846

One hundred parts abrasive grains are coated with 1 part of a resin solvent, e.g., di-butylphthalate, and 6-20 parts of finely divided glycerol-phthalic anhydride reaction product are added, the mixture is warmed to 350° F. to make it plastic and passed several times between rollers, covered with a thin film of linseed oil and maintained at 150° F., and, after final sheeting, articles are cut out and hardened for 48 hours at 350° F.

No. 2 British Patent 434,402

Diamond Dust	26	oz.
Graphite	50	oz.
Charcoal	50	oz.
Red Iron Oxide	75	oz.
Phenol Formaldehyde Resin		
sufficient		ond

Hardness Scale for Abrasives

A scale of hardness based on the lapping method is as follows: bort 10, ballas 9.99, carbonado 9.82, boron carbide 9.32, black silicon carbide 9.15, corundum 9.00.

PYROTECHNICS

Fireworks (Pyrotechnics)

The greatest care should be exercised in making fireworks. Carelessness and impurities produce most accidents. Do not mix large amounts of ingredients and do not permit the introduction of dirt, dust or other foreign matter. Do not mix near your stock of raw or finished material. Make sure that all utensils are cleaned directly before use. Slight friction, even that produced by sifting may cause an explosion or fire. All packing or ramming should be done gently and without scratching as the latter may start a reaction just as well as a shock.

Do not allow matches or open flames in the mixing room. Wear rubber soled shoes. Keep the air moist enough to prevent static sparks from being generated

by moving bodies.

All chemicals used should be of best quality and bought from a reliable house in original packages. These should be kept air-tight. For mixing small quantities round brass wire sieves (No. 16-26) are used. In plain mixings the coal is weighed first and put into bottom of a wooden tub; the sieve is put on top and the sulphur and saltpeter sifted through it. Then with bare arms mix the powder in the tub thoroughly. Place sieve on another tub and sift from first tub a scoopful at a time. Mix with hands again and sift back again into first tub.

In "colored" mixings each ingredient should be sifted separately the first time except the shellac, coal, etc., which is put in bottom of tub. Never throw the chlorate on the sieve with dextrin or other organic material. Beware of hitting the sieve with finger nails or metal-

lic objects.

Sparklers

F	ormula No. 1	No. 2
Lampblack	36	— lb.
Powdered Charcos	al —	25 lb.
Steel Filings	30	50 lb.
Aluminum Powde	er 15	— lb.
Gum Arabic	6	5 lb.
Saltpeter	5	15 lb.
Sulphur	2	6 lb.

The gum arabic is worked up with water into the consistency of mucilage, the other items except the steel filings are stirred in. The steel filing lightly coated with paraffin is finally added. Then work the mixture up to the consistency of porridge.

Pin Wheels

Formula	No. 1	No. 2	No. 3
Meal Powder		10	8 lb.
Fine Grain Powe	ler 8	5	8 lb.
Aluminum			3 lb.
Saltpeter	14	4	16 lb.
Steel Filings	6	6	— lb.
Sulphur	4	1	3 lb.
Charcoal	3	1	8 lb.

Pyrotechnic Fountains

Meal Powder	5	lb.
Granular Saltpeter	3	lb.
Sulphur	1	lb.
Coarse Charcoal	1	lb.
FF Rifle Powder	$\frac{3}{4}$	lb.

Flower Pots

Saltpeter	10	lb.
Sulphur	6	lb.
Lampblack	3	lb.
FFF Rifle Powder	6	lb.

Gerbs

	Formula	No. 1	No. 2
Meal Powder		6	4 lb.
Saltpeter		2	— lb.
Sulphur		1	— lb.
Charcoal		1	1 lb.
Steel Filings		1	2 lb.

Serpents or "Nigger" Chasers

*	Ų.		
	Formula	No. 1	No. 2
Meal Powder		3	3 lb.
Saltpeter		2	5 lb.
Sulphur		1	1 lb.
Mixed Coal		11/2	3/4 lb.
FFF Grain P	owder	4	3 lb

Saltpeter Ammonium Dextrin	Snake Nests Bichromate	1 lb. 2 lb. 1 lb.

Saltpeter

Charcoal

Sulphur

Table Rocket Formula No. 1 No. 2 5 lb. 7 Meal Powder 12 lb. 3 lb. 2 3 lb. Steel Filings

— 1b.

Roman Candles

Powdered Saltpeter	18 lb.
Fine Powdered Charcoal	11 lb.
Flowers of Sulphur	6 lb.
Dextrin	1 lb.
Water	1 gal.

After all the ingredients are well mixed and sifted 3 times, add the water and mix again until the whole lot is evenly dampened.

Rocket and Candle Match

Into a small tub put about a gal. of starch, well boiled, and stir into it about 5 lb. of a thoroughly mixed composition made of

Saltpeter	16	lb.
Fine Charcoal	5	lb.
Sulphur	21/2	lb.

Soak in this, cotton wick of about 5 strands until nearly all the composition is absorbed but about 1/2 in. should still cover the cotton in the tub.

Cascades

Formula	No. 1	No. 2
Granulated Saltpeter	18	16 lb.
Mixed Charcoal	4	4 lb.
Sulphur	3	3 lb.
Iron Borings	6	7 lb.

Smoke Pot

Strontium Nitrate	10 lb.
Sulphur	6 lb.
Whiting (Chalk)	4 lb.
Fine Charcoal	¾ lb.
Dextrin	¾ lb.
or	
Saltpeter	4 lb.
Lampblack	1 lb.
Charcoal	1 lb.
Red Arsenic	1 lb.
Rogin	1 lh

Gold and Silver Rain (Cut Stars)

Formul	a No. 1	${\rm No.}2$	No. 3
Meal Powder	16		4 lb.
Saltpeter	10	1	1 lb.
Sulphur	10	1	— lb.
Fine Charcoal	4	1	2 lb.
Lampblack	2		— lb.
Red Arsenic	1		— lb.
Shellac	1		— lb.
Dextrin	1		— lb.
Lead Nitrate		3	- lb.

Japanese Stars

Formula	No. 1	No. 2
Lampblack	12	6 oz.
Potassium Chlorate	8	4 oz.
Saltpeter	1	- oz.
Water	18	9 oz.
Alcohol	4	2 oz.
Dextrin	1	oz.
Gum Arabic		1/2 oz.

Mix the dextrin and saltpeter together and add sufficient water to make a gummy liquid. Boil the balance of the water and add the potassium chlorate to it. Put the lampblack in a large pan and pour the alcohol over it working it in as well as possible. Then add the potassium chlorate in the hot water and stir with stick until cool enough for the hands and lastly add the dextrin and saltpeter.

In Formula No. 2 the potash and lampblack are sifted together several times; add alcohol; then water in which gum has been dissolved and proceed as in Formula No. 1.

White Stars

Formula	No. 1	No. 2
Saltpeter	50	54 lb.
Sulphur	15	15 lb.
Red Arsenic	15	9 lb.
Dextrin	3	3 lb.
Black Antimony	******	15 lb.
Red Lead		6 lb.
Shellac		1 lb.

Red Stars

Formula	No. 1	$^{\circ}Nc$. 2	
Potassium Chlorate	6	24	lb.	
Shellac or Red Gum	1	3	lb.	
Fine Charcoal	2	4	lb.	
Strontium Carbonate		4	lb.	
Strontium Nitrate	6		lb.	
Dextrin	1/2	11/2	lb.	

Blue Stars Potassium Chlorate 24 lb. Paris Green 9 lb. Barium Nitrate 8 lb. Shellac 5 lb. Dextrin 1½ lb.	Each ingredient should be sifted separately and then mixed in a tub with the fingers, preferably gloved, being careful not to scratch the bottom of tub with the nails.
Chinese Fire Crackers Formula No. 1 No. 2 Saltpeter 50 45 lb. Sulphur 25 18 lb. Charcoal 25 25 lb. Potassium Chlorate — 8 lb. Sand — 4 lb.	Japanese or Cap Torpedoes Formula No. 1 Potassium Chlorate 5 oz. Sulphur 14 oz. Chalk 14 oz. No. 2 Amorphous Phosphorus 2 oz. Sift separately the ingredients of No.
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	1, mix thoroughly and moisten in a bowl with water until of the consistency of porridge. In another bowl moisten the 2 oz. of amorphous phosphorus, to the same consistency. Then stir the phosphorus into the bowl containing the other ingredients with a spoon.
Cannon Cracker Composition Formula No. 1 No. 2 No. 3 Potassium Chlorate 60 6 6 lb. Washed Sulphur 23 3 2 lb. Sulphuret Antimony 5 — lb. Metallic Antimony — 1 lb. Charcoal — 1 — lb. Saltpeter 12 — lb.	White Fire Formula No. 1 No. 2 No. 3 No. 4 Saltpeter 3 12 8 7 lb. Sulphur 1 2 2 2 lb. Metallic Antimony 1 — — lb. Sulphide of Antimony 1 1 — — lb. Realgar — 1 1½ lb.
Red Formula Nitrate of Strontia. Potassium Chlorate Shellac Sheel-lac or Kauri Gum Charcoal Dextrin Fine Sawdust Rosin Lampblack	$\begin{array}{cccccccccccccccccccccccccccccccccccc$

Blue Fire Formula, No. 1 No. 2

In order to make tableau fires more bulky, one to two parts of fine sawdust may be mixed with any of the above formulas without materially affecting the

Chlorate of Potash

Paris Green

Barium Nitrate

Calomel

Sal Ammoniac

color. It should also be borne in mind that paris green is very poisonous and a handkerchief should be tied over the nose if it has to be handled much.

No. 3

1

8

6

1

No. 4

12 1b.

lb.

lb. lb.

lb.

lb.

1½ lb.

	Green Fir	e		
Form	ula No. 1	No. 2	No. 3	
Barium Nitrat	te 8	9	4	1b.
Potassium Chl	orate 4	3	2	lb.
Shellac		1	$1\frac{1}{2}$	lb.
Sheel-lac (She	llac			
Substitute)	2			lb.
Dextrin		1/16		lb.
Fine Sawdust		1/2		lb.
Sal Ammoniac	1			lb.
_				

Yellow Fire	
Barium Nitrate Sodium Oxalate Sulphur Sheel-lac	36 lb. 6 lb. 3 lb. 5 lb.
Red Lances Formula No. 1	No. 2
Potassium Chlorate 16 Strontium Nitrate 3 Strontium Carbonate — Shellac 3 Lampblack ½	16 lb. - lb. 3 lb. 2 lb. 1 lb.

Green Lances

Formula N	o. 1	No. 2	No. 3	No. 4
Potassium Chlorate	7	16	16	— lb.
Barium Nitrate	. 7	4	6	— lb.
Barium Chlorate				6 lb.
Shellac	2	4	3	1 lb.
Calomel		3	3	2 lb.
Lampblack		1/8		— lb.
Dextrin			1	— lb.
Pieric Acid			1	1 lb.

White Lances

Formula	No. 1	No. 2	No. 3	No. 4
Saltpeter	9	14	5	8 lb.
Sulphur	1	4	2	2 lb.
Antimony Sulphio	le 2			— lb.
Antimony Metal				
Powder		3	1	— lb.
Meal Powder			1	— lb.
Red Arsenic	,			1 lb.

Magnesium Torches

a.	Shellac	120 g.
	Resin	120 g.
	Barium Nitrate, Dry	840 g.
b.	Magnesium Powder	25-40 g.

Mix the ground a with b, and fill into zinc-tubes (thin walls) having a wooden handle, which closes the tube below.

Parade Torches

Strontium Nitrate		40	lb.
Potassium Chlorate		. 8	lb.
Red Sheel-lac		7	lb.

Railway Fuses

No. 1	No. 2	No. 3	No. 4
e 48	16	18	16 lb.
12	4	7	4 lb.
5	2	2	5 lb.
4	1	1/2	1 lb.
10	3	2	— lb.
		1/2	— lb.
	e 48 12 5 4	e 48 16 12 4 5 2 4 1	$ \begin{array}{cccccccccccccccccccccccccccccccccccc$

Ship Distress Signals

		0		
Potassium	Chlorate		5	lb.
Strontium	Carbonate		$1\frac{1}{2}$	lb.
Shellac			1	lb.
Dextrin			11/2	lb.

Miracle Candles

	THE COLO COMMUNICO	
a.	Iron, Powdered	25 g.
b.	Barium Nitrate	52 g.
c.	Aluminum Powder	8 g.
d.	Starch, Wheat	15 g.

Right size of the iron grains is most important, b and c should be finely powdered.

Should be produced in summer for quicker and more economical drying. Mixture must be perfect, pack in airtight drums.

Put into an enameled container (best way using 1 kg. mass), make a little hole in the center of the powder, pour in it the least possible amount of boiling water (100 g. for 1 kg. powder), and stir the whole thoroughly. The right point of pastification and right amount of water is reached when the paste is not too friable or too sticky and forms a concrete non-sticky mass.

This mass is put on wires (2 g. per wire), and dried.

Orange Smoke U. S. Patent 1,975,785

A pyrotechnic composition for producing orange smoke, comprises lead peroxide 50 parts, potassium bichromate 35 parts, and magnesium 15 parts.

Brown Smoke

U. S. Patent 1,975,099

A pyrotechnic composition for producing brown smoke, comprises copper oxide 50 parts, lead peroxide 35 parts, and magnesium 15 parts.

Pyrotechnical Device U. S. Patent 1,936,221

A firework of the "sparkler" type consists of an iron rod coated at one end with a plastic mixture of barium nitrate

S5, strontium carbonate 60, sodium aluminum fluoride 40, potassium chlorate 225, dextrin 30, and shellac 55 all parts by weight in which are embedded granules of magnesium-copper or magnesium-aluminum alloy.

Explosives

Formula No. 1 British Patent 408,260

Explosives consist of alpha-trinitrotoluene 10-30, o-nitrotoluene 5-10, ground coconut fiber or charcoal 1-5, paraffin or other suitable wax 3-6, aluminum, graded 50-mesh, 10-24, finely powdered aluminum 1-4, and barium nitrate, or other nitrate, 70-21 parts,

No. 2

British Patent 412,583

A nitrated mixture of glycerol and glycol 15, ammonium nitrate 8.5, sodium nitrate 12.0, plant fiber 6, sodium chloride 58 and ammonium orthophosphate 0.5%, has a density of 1.1 g. per cc. and gives a ballistic pendulum swing of 1.08 in., the volumetric power factor being 1.19.

No. 3

British Patent 435,588

Ammonium Nitrate 90 lb. Aluminum Powder 6½ lb. Manganese Dioxide Powder 3½ lb.

Slow-Burning Explosives British Patent 423,040

Examples of slow-burning explosives are (1) potassium nitrate 75 or sodium nitrate 73, charcoal 15 or 17 and sulphur 10, (2) sodium nitrate 44, ammonium nitrate 34 and charcoal 22%. The explosive may be granular or compressed in pellets and may contain small quantities of cooling salts and boric acid or borates.

Explosive Priming Mixture British Patent 432,096

A suitable composition is the potassium salt 16, basic lead salt of trinitroresorcinol 15, barium nitrate 40, and antimony sulphite 29%.

Priming Charge Canadian Patent 348,291

A solution containing potassium nitrate 30, barium nitrate 20, and water

100 parts is crystallized at 50° C. to give a double salt, which when used in priming charges leaves substantially no corrosive residues nor fused masses in the barrels of firearms; e.g., a priming charge consists of mercury fulminate 20–45, potassium barium nitrate 30–60, lead thiocyanate 10–40% by weight.

Priming Composition German Patent 614,712

The composition contains zirconium powder in addition to the usual constituents. Thus, the composition may contain zirconium powder 10, barium nitrate 40, mercuric fulminate 25 and antimony trisulphide 25%.

Flash Composition U. S. Patent 1,964,077

A suitable mixture contains perchlorate 20, potassium chlorate 39.5, silver nitrate 39.5, and nitrocotton 1.0%.

Flashlight Cartridges British Patent 419,658

A cartridge is charged with a powder mixture consisting of magnesium 700-900, sulphur 10-18, potassium permanganate or potassium chlorate 100-140, potassium nitrate 70-85, magnesium oxide 100-160 and charcoal 10-13 parts.

Black Powder Canadian Patent 348,641

The addition of 0.1-5.0% by weight of stearic acid retards the burning speed of black powder. E.g., a black blasting powder contains sodium nitrate 72.0, sulphur 10.0, charcoal 17.7 and stearic acid 0.3%.

Fuse Powder French Patent 783,249

A powder of long combustion is made by dissolving niter 5, pulverized sulphur 4 and wood charcoal 3.5 parts in pure alcohol to form a thick mass which is well mixed and dried.

Gelatin Dynamite Canadian Patent 352,763

The following percentage compositions are specified:

Formula No. 1

Nitroglycerin 4 Dinitrotoluene

Nitrocotton Sodium Nitrate Expanded Cereal Product Starch Chalk	1.3 36.1 9 2.7 0.9
No. 2	
Nitroglycerin Dinitrotoluene Nitrocotton Sodium Nitrate Ammonium Nitrate Expanded Cereal Starch Chalk	60 3.5 2.3 2.2 24 6 1
No. 3	
2.0.0	30
Nitroglycerin Dinitrotoluene Nitrocotton Sodium Nitrate Ammonium Chloride	2 0.7 44.8 15
Expanded Cereal Product Starch Chalk No. 4	$\frac{2}{4.5}$
Nitroelyaarin	22
Nitroglycerin Dinitrotoluene Nitrocotton Sodium Nitrate Ammonium Nitrate Expanded Cereal Product Chalk	1.5 0.2 9 60 6.9 0.4

Detonators

French Patent 781,646

A composition which is fired directly by the passage of an electric current comprises a mixture of finely divided zirconium and a nitrophenol salt of lead, e.g., zirconium 70 and lead mononitroresorcinate 30 parts in sufficient amount of a 5% solution of nitrocellulose in amyl acetate to make a creamy mixture.

Percussion Detonator U. S. Patent 1,975,679

A percussion detonating composition consists of phosphorus sesquisulphide 30 g., gum arabic 115 g., magnesium carbonate 20 g., calcium carbonate 5 g., potassium chlorate 80 g., iron sesquioxide (red ochre) 40 g.

Waterproofing for Blasting Fuses (Non-Staining)

U. S. Patent 1,968,907

Petrolatum	25-90	lb.
Ester Gum	5-75	lb.
Paraffin Wax	5– 50	lb.

RUBBER, RESINS, WAXES, PLASTICS

Caoutchouc (Rubber) Synthetic
Acetylene is absorbed by a mixture of
Cuprous Chloride 1000 g.
Ammonium Chloride 400 g.
Copper 100 g

Copper 100 g.
Hydrochloric Acid
Concentrate 30 g.
Water 425 g.

at 40-50° C. The saturation is reached, when 50 g. of acetylene have been absorbed (3 hours). The mixture is kept at ordinary temperature during 24 hours, then distilled on an oil bath. The distillate contains 33% of

 ${\rm CH_2} = {\rm CH} - {\rm C} \equiv {\rm CH},$ and 67% of superior condensation products, among which has been found

 $CH_2 = CH - C = C - CH = CH_2$, and C_8H_8 .

In the same process, the yield in $CH_2 = CH - C = CH$

falls, when the period between saturation and distillation is increased to 140 hours.

A 70% yield is obtained when running the absorption at 80° C., and collecting the gas of reaction into two receivers, the first chilled in ice, the second in carbon dioxide snow. The liquid in the second receiver contains:

 $CH_2 = CH - C = CH$ CH = CH $CH_3 - CHO$

The chloroprene is obtained with an 80% yield, agitating

 $\begin{array}{c} {\rm CH_2 = CH - C \equiv CH \ with} \\ {\rm Hydrochloric \ Acid,} \\ {\rm Concentrate} \\ {\rm Cuprous \ Chloride} \\ \end{array} \begin{array}{c} 70 \ {\rm g.} \\ {\rm Cuprous \ Chloride} \\ \end{array}$

Ammonium Chloride for 3 hours at room temperature.

Rubber Master-Batch U. S. Patent 1,942,853

Substantially unmasticated crude rubber (1 lb.) is superficially treated with ½-3 lb. of a softener, e.g., mineral oil, so that the latter is absorbed. This procedure obviates the difficulties of incor-

poration of liquid softeners in the usual manner, and the soft, non-tacky product is very easily mixed with other compounding ingredients.

Porous, Fibrous Rubber Compositions British Patent 409,294

A porous, non-waterproof, fibrous, feltlike material is prepared by admixture of rubber with finely comminuted (not powder) fibers of wool and hair in proportions of not more than 50% rubber and not less than 50% fibers together with an amount of non-liquid expanding agent, e.g., ammonium carbonate, sodium carbonate, sodium potassium carbonate, sodium bicarbonate, ammonium bicarbonate that will expand the mass 2-6 times. Vulcanization and coloring agents and softeners may be added. In an example sulphur 7.5, zinc oxide 6, ferric oxide 2, stearic acid 4, ultra-accelerator 1 oz. and comminuted wool 22.5 lb. are added to 15 lb. softened rubber. When cool ammonium bicarbonate is added and the product calendered into sheets.

Rubber Fibers German Patent 614,615

Rubber fibers are formed by introducing a coagulating agent through nozzles into rubber latex. Thus, a 60% solution of acetic acid is fed into a rubber latex mixture of rubber 92.5, sulphur 2.5, zinc oxide 2.5, anti-oxidation agent 1.0, accelerator 0.5 and ammonium oleate 1%, through 0.42 mm. nozzles, the fiber being removed at 600-760 cc. per minute.

Chlorinated Rubber British Patent 410,249

A solution of unvulcanized (artificial or reclaimed) rubber, gutta-percha or balata, with or without factice, admixed with 5-20% uncombined sulphur is chlorinated to yield a thermoplastic mass suitable for the manufacture of films, varnishes or moldable compositions,

4 g.

the chlorination being continued until the gel which forms is entirely redissolved. Metallic halides, oils, turpentine, chlorinated naphthalenes, tritolyl phosphate, organic esters, ethereal oils, cellulose plastic softeners, synthetic resins or varnishes may be added before, during or after chlorination. In an example, 10 g. masticated crepe in 200 cc. carbon tetrachloride is mixed with 1 g. sulphur and heated with chlorine until the gel formed redissolves to form a mobile liquid and the product is precipitated by adding 100 cc. alcohol and washed in boiling water to give a white mass containing 32% chlorine, soluble in acetone and benzol to yield a transparent, colorless film moldable at 130° C. If the chlorination is stopped before resolution, the gel which rises to the surface being removed, washed with solvent, treated with boiling water and dried, the product will be a semitransparent, hard, tough, substance moldable at 130-140°.

De-Polymerization of Rubber German Patent 599,405

Rubber can be de-polymerized to give 40-60% solutions by treatment in suspension or solution with 10% of its weight of 53% nitric acid. A paste is first prepared by stirring 10 kg. rubber in 90 kg. benzol, whereupon 1 kg. of the 53% nitric acid is stirred in and the de-polymerization interrupted at the desired stage by neutralization with ½ kg. barium carbonate.

The de-polymerized rubber solution is decanted off and concentrated if necessary by evaporation. Coatings of this form of rubber are somewhat tacky but this defect can be remedied by a partial re-polymerization (immediately after the neutralization stage) with antimony trichloride or phthalic acid in alcoholic solution.

Cork-Rubber Composition British Patent 425,699

Rubber	100	lb.
Cork	100	lb.
Sulphur	3	lb.
Zinc Oxide	5	lb.
Stearic Acid	2	lb.
Mercaptobenzothiazole	0.5	lb.
Zinc Isopropylxanthate	Piperi-	
dine-1-Carbothionolate	0.5	lb.
Paraffin	5	lb.
Nonox S	1	lb.
Lithopone	25	lb.
Chromium Oxide, Green		lb.

Cork Composition Canadian Patent 348,152

A mixture of phenol 13, paraformaldehyde 8 and diethylene glycol 30 parts by weight is heated to 210° F., 6.4 parts by weight of a 16% solution of caustic soda is added as a catalyst, and the heating is continued at about 210° F. until a sample of the liquid taken off will set in 10 minutes in boiling water. The product is immediately mixed with ground cork in the proportion of 80 lb. of the liquid and 150 lb. of cork particles. The treated cork is placed in a mold at about 300° F., where the reaction is completed and the comminuted cork is agglomerated into a cohesive mass of the desired shape.

Coating for Rubber Goods British Patent 427,228

Latex	100 lb.
Glue	1-5 lb.
Barytes	100 lb.
Titanium Dioxide	50 lb.
Rosin Oil	10 lb.
Casein	5-20 lb.
Sulphonated Castor Oil	5 lb.
Ammonia (28%)	8 lb.
Formaldehyde	10 lb.
Color	to suit

Water sufficient to give a final concentration of total solids of 45-50%.

Thermoplastic Hornlike Rubber German Patent 615,050

Treat rubber with 70% hydrofluoric acid for 24 to 48 hours.

Rubber Curing Solvents Formula No. 1

Carbon Bisulphide	50 gal.
Petroleum Naphtha (140-220° F.)	50 gal.
Sulphuryl Chloride	1 gal.
No. 2	
Carbon Tetrachloride	50 gal.
Petroleum Naphtha	
(140-220° F.)	50 gal.
Sulphuryl Chloride	1 gal.

Fire-Resistant Rubber U. S. Patent 1,966,271

Formula No. 1

A solution of 100 parts ammonium chloride, 6 parts ethylene glycol, and 3 parts glue in 300 parts of water is added to 3 parts of an antioxidant comprising a mixture of the condensation products of acetaldehyde with a.and. \(\text{g-naphthyl-amines} \), the antioxidant being wetted with a little alcohol. Sponge rubber is soaked in this solution and the excess squeezed out until the "wet" gain in weight is 120% on the dry weight. It is dried in a current of warm air. Sponge rubber treated in this way will not burn unless held continually in a flame, and on withdrawal from the heating flame, the sponge at once ceases to burn.

No. 2

Five parts casein are dissolved in ammonia solution, the bulk made up to 300 parts and 100 parts ammonium chloride dissolved in it. The sponge rubber is soaked in this solution and the excess squeezed out so as to leave in the sponge a quantity of solution equivalent to 120% of the weight of the dry sponge rubber. This is then dried in a current of warm air. The degree of fire-resistance can be adjusted by alterations in the proportion of ammonium chloride present.

No. 3

Excess selenium is boiled for 30 minutes with 20% ammonium sulphite solution and the solution obtained is filtered through glass wool. The sponge rubber is soaked in this solution and squeezed out until the increase in weight is 55% of the dry weight, and dried in a current of warm air. The selenium is slowly deposited spontaneously by exposure.

No. 4

Sponge rubber impregnated as in the preceding case with a solution of selenium in ammonium sulphite is exposed to an atmosphere of sulphur dioxide for liberation of the selenium. Alternatively finely powdered selenium is rubbed on to the surface and into the surface pores of sponge rubber so that some is permanently retained. The extent of fireresistance depends upon the quantity of selenium retained.

No. 5

Sponge rubber soaked in a 20% solution of ammonium silico-fluoride resqueezed out until the gain in weight is 150% of the dry weight. It is dried in a current of warm air. Sponge rubber treated in this way will not burn unless heated continually in a flame, and on withdrawal from the heating flame at once ceases to burn.

No. 6

Thirty parts of finely powdered ammonium chloride are stirred into 100 parts of a 1% solution of rubber in benzene, and the suspension obtained is painted on to the surface of the sponge rubber. The solvent is allowed to evaporate and the surface is dusted with French chalk. The fire-resistance of the sponge rubber is markedly improved. A similar suspension of ammonium silicofluoride in a benzene solution of rubber has a like effect.

It is to be understood that the quantity of fire-preventing agent remaining in the pores is not sufficient to fill the latter in any case; that the porous structure is not changed and that the agent is deposited as a superficial coating on the inner sur-

faces of the pores.

Fireproofing Rubber British Patent 432,551

Five to fifteen per cent of any of the following is incorporated in the rubber:

Triphenyl Phosphate Tricresyl Phosphate Triphenyl Borate

Rubber Calender Liner

The handling of miles of calendered sheet involves either efficient dusting methods, to permit rolling up of the sheet without risk of adhesion, or alternatively a good non-adhesive cloth that can be rolled up with the rubber. Where the sheet has subsequently to be cut into shapes and built up, its tackiness is important, so that dusting becomes out of the question in industries such as tire and footwear manufacture.

nd rootwear manuracture.		
Gelatin	75	lb.
Glycerin, Commercial	85	lb.
Talc		lb.
Dye, Color to Suit	10	lb.
Water	800	lb.

The cotton is treated with this mixture on both surfaces and dried. It is then hardened by passing through a bath of 10% formaldehyde solution, dried, and pressed on a calender.

One thousand square meters of cotton sheet can be covered with 37.5 kg. gelatin, 42.5 kg. glycerin, 15 kg. tale, 0.5 kg. dye, and 25 kg. formaldehyde.

Rubber Mold Lubricant

Sodium Hyposulphite	280	g.
Sugar	70	g.
Magnesium Sulphate Crystals	30	g.
Glycerin	15	g.

Hexamethylene Tetramine
Phenol
Sodium salt of the sulphuric
acid derivative of the reaction product of normal
butyl alcohol and a mixture of approximately 85%
ortho hydroxy diphenyl and
substantially 15% para

hydroxy diphenyl

The composition thus prepared is added to substantially 20-30 times its weight of water. When applied on the surface of molds and press plates, which contact with rubber or other material to be vulcanized or molded, the film produced is markedly tough and resists rubbing off when the rubber or other material is pressed into the mold.

Lubricant for Vulcanizing Molds
Sodium Hyposulphite 3 lb.
Ammonium Carbonate 1 lb.
Water 97 lb.

Non-Adhesive Mold Liner

To a mixture of casein 45, glycerol 45, and kaolin 10 parts, add water to the required consistency. Apply 2-3 times on both sides of the cotton material. Dry 1-1½ hours. Then treat with formaldehyde. The total time is 6-7 hours.

Aqueous Latex Dispersions for Artificial Leather

A mixture composed of "smoked sheets" (rubber) 100, gasoline 200, oleic acid 8, 25% ammonium hydroxide solution 20, casein 20, sulphur 8, zinc oxide 10, "Kaptax" 2, thiuram 1 part and water in accordance with requirements, produces stable emulsions when diluted with up to 50 volumes of water. A leather substitute of good physical and mechanical properties is obtained from rubber 100, rosin 19, oleic acid 5, wheat flour 15, glue 5, kaolin 10 and sulphur 5 parts.

Rubber Films and Threads

A hydrochloric additive product (1) is prepared, in 100% yield, by treating a 2% benzol solution of rubber with hydrochloric acid at 16.5-19° C.; after 15 hours the product is separated by precipition with alcohol. Glossy, transparent films may be prepared by spreading chloroform solution of (1) on a glass plate and allowing it to evaporate at 45-50° C. The films adhere to metals and may be

dyed; they may be combined with plasticizers, which reduce the strength but increase the extensibility. Threads may be prepared by dry spinning from a 7% chloroform solution of (1). The material is not readily combustible, and is but little acted on by hydrochloric acid (concentrated and 2N), potassium hydroxide (20% and 2N), soap solutions, or 4N-sulphuric acid. Decomposition is effected by treatment with concentrated sulphuric or nitric acid or by prolonged heating at 55-60°.

Hard Rubber Coating U. S. Patent 2,023,582

A method of applying a hard rubber coating to articles comprises mixing substantially 500 g. of smoked sheet rubber, 180 g. of sulphur, 2½ g. of diphenylguanidine, and 2½ g. of mercaptobenzothiazole, dissolving the mixture in substantially 2500 g. of benzine, applying the solution to an article to form a coating and vulcanizing the coating.

Wire Insulation Compound

The following formula provides an insulating compound capable of extremely rapid vulcanization and yet one which, when mixed and applied in accordance with the process defined, does not vulcanize during the extruding operation.

Smoked Sheet Rubber	22	g.
Reclaimed Rubber (Boot	7.0	
and Shoe)	10	\mathbf{g} .
Reclaimed Rubber (Whole		
Tire)	10	g.
Mineral Rubber	5	g.
Whiting	44.7	g.
Zinc Oxide	2.5	g.
Antioxidant	1.5	g.
Sulphur	1	g.
Softener (Pine Tar Oil)	3	g.
Ultra-Accelerator	0.3	

This stock is adapted for continuous vulcanizing carried on at a high rate of speed. For example in coating No. 17 Brown and Sharpe gage drop wire with a coating 3/4 in. thick satisfactory results are obtained when the speed of travel is from 400 to 500 ft. per minute when using a vulcanizing chamber 100 ft. long. The corresponding vulcanizing periods for these speeds would be 12 to 15 seconds.

Molded Brake-Lining U. S. Patent 1,963,511

and allowing it to evaporate at 45-50° C.

The films adhere to metals and may be or water comprises asbestos 29.7, carbon

black 7.7, barium sulphate 12.8, lead oxide 2.0, rubber 33.2, sulphur 4.6, and an aqueous suspension of a phenolic or other infusible resin 10 volume per cent.

Electrical Insulation for Cables

Satisfactory insulation is achieved by coating the cable with a vulcanized mixture of synthetic rubber 15, filler (kaolin, chalk) 40, and asphalt 45%.

Artificial Leather (Gralek)

A fabric is coated with

10% Rubber Solution in
Gasoline 87½ lb.
Zinc Oxide 8 lb.
Sulphur 3 lb.
Thiuram 1½ lb.

and then with a mixture of

Rubber 100 lb. Leather Dust 150-200 1b. Zinc Oxide 8 lb. 2 Sulphur Ib. 20 lb. Lampblack Accelerator 11/2 lb. Pine Tar 5 lb. Gasoline 1000-1200 lb.

The final coating consists of dry casein pigment, formalin 6% of dry pigment and alizarin oil 10%. Finally vulcanize and varnish.

Transmission Belt Dressing U. S. Patent 2,001,582

Neatsfoot Oil 1 lb. Rubber 13 lb.

Printing Blanket Patented

Formula No. 1

A flexible and resilient printer blanket having a smooth surface which is resistant to oils and repellent to inks is made by applying a chlorinated rubber coating over the ordinary printer blanket. The coating varnish comprises chlorinated rubber (20 to 40), benzene (10 to 75), a plasticizer (3 to 10). Another varnish may contain chlorinated rubber (30), xylene (25), tricresyl phosphate (5); while in the other example there are combined chlorinated rubber (30), benzene (30), dibutyl phthalate (6). When it is desired to use pigments or dyes in the varnish it is preferred that the pigment, such as carbon black, is first mixed with the plasticizer and then incorporated in the chlorinated rubber solution.

No. 2 British Patent 423,556

In a printers' blanket of the type comprising a fibrous base and an outer coating of, or containing rubber, which is surfaced or ground in the usual manner, the outer coating is obtained directly from an aqueous dispersion of, or containing, rubber and is vulcanized to the base. The fibrous base may comprise a plurality of superposed layers of fabric material, e.g., felt, that are bonded together by a rubber cement or latex adhesive, containing vulcanizing ingredients, so that on subsequent vulcanization of the rubber coating the adhesive is also vulcanized. A preferred latex composition comprises rubber (as latex of 65.7% solids content) 100, formalin 4.65, water 79.75, potassium hydroxide 0.90, antimony sulphide 20, sulphur 5.8, whiting 75, ferric oxide 12, zinc oxide 2, sodium isopropyl naphthalene sulphonate 0.975. glue 0.375, heptaldehyde aniline condensate 1.5, acetone diphenylamine condensate 0.75 and solvent naphtha 1.5 parts; the coatings are dried at 90° C. and vulcanized at 135° C.

Mending Rubber Goods

Apply to the surface of the object a thin solution of rubber in benzol such as is used for sticking patches to auto tubes and allow a few minutes to evaporate solvent. Apply a generous coating of latex rubber and allow to stand a few hours. Can be used for mending auto tops, cuts in tires, hot water bottles, etc.

Rubber Packing Rings for Grooved Cans In grooved containers with rubber packing rings the caps are set in place and the rings heated to 150-180° F. under pressure for about 1 sec. The formulation of the rubber ring is of importance for the proper speed of melting and the proper degree of hardness. A typical formula is (in percentages by wt.): rubber 14.10, balata 4.70, heavy spar 55.56 and chalk 25.64.

Puncture Proofing Tire Tube

A self-healing inner tube structurally designed to prevent deflation after puncturing is secured by lining the tube during its manufacture with a tread ply of rubber of special softening composition. The following formula gives satisfactory results:

Phosphoric Acid Clay

2 lb. 134 lb.

Rosin Oil 3 lb. Rubber 931/4 lb.

The particular softening agent used is ortho-phosphoric acid of 85% strength. The clay serves as a vehicle for the phosphoric acid. The clay and acid are mixed together before being added to the other ingredients. The rosin oil serves as a softener and tack producer. The ingredients are mixed on a rubber mill in the usual manner and may be calendered and slit into strips. In the construction of an inner tube by the pole or flat drum method one of these strips is used as a lining for that half of the tube toward The application of heat to the tread. the tube results in vulcanization of the body structure, but the special stock layer, due to the presence of the chemical agent and absence of sulphur, accelerator, or other vulcanizing ingredients in its composition, does not vulcanize. On the contrary it becomes extremely plastic, almost viscous in form, and interiorly is very sticky. Although the non-tacky layer in the tube causes the surface of the special stock layer to be somewhat less sticky so that it will not adhere to the opposite wall of the tube should it come in contact therewith, it is preferable that the finished tube be kept in lightly inflated condition. In the event of puncturing by a nail the sticky layer adheres to the nail so that when the nail is withdrawn, it draws back some of the sticky stock with it so as completely to seal the hole through the body structure.

Puncture Proofing Tires German Patent 589,394

Use is made of mixtures of latex with animal, vegetable or mineral oils. A typical mixture contains ammoniacal latex 40, sesame oil 50 and olein 10%. The mixture is introduced through the air valve of the tire, distributes itself over the inner surface and automatically seals any punctures which may develop.

Gas Generating Composition for Rubber Balls

A stable mixture of ingredients from which to prepare pellets for use in inflating hollow balls, etc., follows:

Ammonium Chloride 40 lb. Sodium Nitrite 59 lb. Anhydrous Sodium Carbonate 1 lb.

The main constituents, viz., the ammonium chloride and the sodium nitrite, are commercial materials not completely dried. When maintained at 60° C., this

gas producing mixture decomposes roughly 25 to 30 times more slowly than pellets prepared from dried materials but without sodium carbonate, and over 100 times more slowly than pellets prepared from undried commercial materials, again without sodium carbonate. They undergo no appreciable decomposition at ordinary temperatures, or is their value diminished for inflating rubber balls at 100° C. (212° F.) or over.

Rubber Vulcanization Accelerator U. S. Patent 1,963,084

Turpentine 100 oz. Sulphur 15 oz. Heat at 120–130° C. for 12 hours.

Forms for Molding "Bakelite"

Graphite Powder	4 g.
Clay	4 g
Magnesia	2 g.
Cement	12 g
Asbestos Powder	1 g.
Magnesium Chloride	8 g.
Mold to a paste.	

Dental Thermoplastic Molding Composition

U. S. Patent 2,020,311

Twenty-five parts of rosin are melted or fluxed with 1 to 5 parts of glycerol (depending upon the abietic acid content of the rosin), preferably under a reflux condenser, and from 10 to 25 parts of aluminum stearate added to the mixture while it is still at a relatively high temperature, that is, 250° C. or there-From about 5 to 10 parts of rosin oil are then added, if desired, and after this has been thoroughly incorporated into the body, it is allowed to cool to a temperature of about 150° C., whereupon from 1 to 5 parts of triethanolamine stearate is added. Thereafter, wood flour may be incorporated. Prior to the addition of triethanolamine stearate, the composition, although elastic, is extremely sticky and gummy and unsuited for dental purposes.

Dental Impression Jelly

	-		
Agar-Agar		14	g.
Water		100	g.
Glycerin			minims
Kaolin		12	ø.

Dissolve agar-agar in water by heating in a pressure cooker for 1½ hours. Then stir in other ingredients.

Plastic Molding Composition U. S. Patent 1,969,146

Phenol Formaldehyde Resin	4	lb.
Charcoal, Powdered	6	lb.
Wood Flour	3	lb.
Pine Tar	1/2	lb.

Capsule Composition (Cheap)

Potassium Silicate	
(30-33° Bé.)	70 g.
Water-Soluble Dye	2 g.
Water	28 g.

Capsule Composition

Gelatin	27 g. 42.7 g.
Water	_
Allow to swell over night gently with stirring until uni	and warm form.
Glycerin (28° Bé.)	10 g.
Water-Soluble Dye	2 g.
Water	18 g.
Preservative	

Manufacture of Casein

"Rennet Casein" suitable for making Galalith and similar plastics is best obtained as follows: To fresh skim milk at 35° C. add sufficient rennet to effect coagulation in 15–20 minutes; stir 5–10 minutes and warm to 65° C. at the rate of 1° per minute; decant twice with water at 25°; drain and press out as much water as possible, disintegrate the press cake and dry at 43–45° C.

Plastic Composition French Patent 781,749

A composition for making pipes contains asbestos 85, fluid resin 15, lithopone 0.15, muldrite 1500 and cellulose 2200 kg. or vegetable fibers 85, resin 10 and rubber or latex or bitumen 5 kg.

Plastic Display Composition

Compositions based upon pigmented linseed oil, castor oil, and a non-alkaline thickening agent such as corn starch, have recently been suggested as constructional material for display work. They are also eminently suitable for coating theatrical drop curtains and the like. They can be produced in various colors, and of a consistency permitting easy stencilling.

For a yellow compound, 16 oz. of a paste pigment in the ratio of 6 lb. white

lead to 4 lb. chrome yellow are worked up into 80 oz. of spar varnish, 10 oz. boiled linseed oil, 10 oz. Japan drier, and 2 oz. castor oil. Sufficient corn starch is then incorporated to yield a mass with the consistency of thick mortar, which is allowed to mature in the open air for about 12 hours before packing into airtight containers. Castor oil is an essential ingredient, since it assists maintenance of the solids in suspension for a very long period if the containers are air-tight.

When making up a bright red or orange composition in which the pigments accelerate drying, the above formula must be modified to the extent of using more castor oil (3 to 5 oz.), more linseed oil and less spar varnish. On the other hand, the slow-drying black compositions will require a higher proportion of varnish and japan, and as little as ½ to 1 oz. castor oil. This type of composition appears to be suitable for producing numerous figures required in industrial display work, the advantages being maintenance of flexibility and toughness after drying, good adhesion to supports and resistance to chipping.

Modeling Clay Formula No. 1

What is called molding compound by some artists is made by mixing two parts by weight of kaolin or powdered soapstone, which must be bone dry, and one part by weight of wheat flour, stirred into three parts of melted white beeswax (not too hot), and well kneaded before the wax cools. The mass may be colored to suit. A good modeling clay can be made from dry clay, mixed with glycerin instead of water. The mixture must be thoroughly mixed.

	No.	2		
Plastic Clay			46	oz.
Cup Grease			24	oz.
Paraffin Wax			11	OZ.
Rosin Oil			1	oz.

Polishing Plastics

Cast resins polish to a high, permanent luster. Rough cuts are usually ground, using the same type of equipment as required by wood or brass. Sand paper, garnet paper, belts or fine abrasive wheels are used. For most work, a generous supply of water is recommended, when wheels are used, to prevent overheating and to keep the wheel clean.

Surfaces which show tool or grinding marks are given a smooth surface, preparatory to final polishing, by "ashing," in which an ordinary buffing wheel, made of muslin discs, of 12 to 14 in. diameter, is used. Wet pumice, kept in a shallow pan under the wheel so that the buff just touches it, is used as a polishing agent. Often, additional wet pumice, taken from the trough, is applied by hand or trowel above the piece being worked. Polishing is usually done, on larger pieces, by a second wheel, using bar wax or specially prepared polishing compounds. These wheels, usually 12 in. in diameter, operate around 1800 r.p.m. A third, clean dry wheel is used to give a final polish.

Tumble Polishing

For large quantities of small and medium sized pieces, tumbling is often used. Here, barrels of hard wood, lined with leather or heavy felt and operating at about 50 r.p.m. are used. Solutions vary with the article being polished, a common procedure calling for preliminary tumbling in dry pumice, to which wooden shoe pegs or similar agents have been added to provide friction. The pumice is later washed off and a second tumbling follows in damp hard-wood sawdust. Other materials are sometimes used as well as pumice. A final operation consists in tumbling with powdered stearic acid or red oil. In some cases emulsions of carnauba wax are used.

PROPERTIES OF NATURAL RESINS

						Direct
	Per		S	often-		\mathbf{Acid}
	cent	\mathbf{Direct}	Indirect	ing	Melting	Number
	Mois-	Acid	\mathbf{Acid}	Point	Point	\mathbf{After}
Natural Resins	ture	Number	Number	°C.	°C.	Running
Genuine Bold Pontianak	1.5	123	133	108	141	95
DBB Soluble Copal Chips	2.4	139	157	90	119	97
No. 1 Brown Kauri	5.4	57	67	120	152	35
Bold Black Scraped	1.5	20	36	125	164	17
Batu Bold Scraped	3	18	33	132	180	15
Pale Bold E. I. Singapore	0.7	20	37	128	156	9
Hard Dark Amber Congo	0.7	102	123	104	200	78
Congo Gum, Ivory Rescraped	1.8	92	111	91	144	92
Medium Pale Congo	0.4	110	132	85	220	70
Boea Medium Dark	2.9	126	149	115	148	95

Softening Point determined by the capillary tube method.

Melting Point determined by the Mercury Method-Rangaswami, reported in the
Journal of the Oil and Color Chemists Asso., 1930, Vol. 13, Page 287.

CLASSIFICATION OF NATURAL RESINS

- L Low Acid Number Resins, including Damar and East India type.
 - A. Damar Resins—oil soluble—indirect acid number 25-45 M.P. 90-110° C.
 - 1. Batavia
 - 2. Sumatra
 - 3. Pontianak
 - 4. Padang
 - 5. Singapore
 - B. East India Fossil or Semi-fossil Resins—oil soluble—indirect acid number 25-40 M.P. 125-180° C.
 - 1. Batu
 - 2. Hiroe
 - 3. Rasak
 - 4. Macassar East India
 - 5. Bold Black Scraped
 - 6. East Indian Singapore
- II. Resins of High Acid Number originating in the East Indies:
 - A. Pontianak—Fossil resins—oil and spirit soluble—indirect acid number 103-140 M.P. 135-145° C.

B. Manila resins

1. Soft or Menlengket resins-spirit soluble-indirect acid number 135-160 M.P. 110-135° C. Macassar

2. Half hard or Loba resins-spirit soluble-indirect acid number 140-

150 M.P. 115-120° C. Loba and Macassar Loba

3. Hard fossil resins-oil and spirit soluble-indirect acid number 105-120 M.P. 140-155° C. Boea-Loewoe-Pontianak

III. African Fossil or Semi-fossil oil soluble-indirect acid number 110-135 M.P. 140-220° C.:

A. Congo

IV. New Zealand fossil or semi-fossil resins-oil and spirit soluble-indirect acid number 55-70 M.P. 120-160° C.:

A. Kanri

B. Bush Kauri

Melting Points of Synt	hetic Resins
Amberol BS1	99-110° C.
Bakelite BR352	93-104° C.
Bakelite BR2072	80- 91° C.
Beckacite 1101	102-112° C.
Beckacite 1102	102-112° C.
Beckacite 1113	102-112° C.
Akco Resin, Hard	125-130° C.
Amberol F7	118-125° C.
Amberol 226	117-133° C.
Amberol 801	117-133° C.
Beckacite 1112	110–125° C.
Lewisol No. 1	120–125° C.
Paranol, Hard	115° C.
Paranol, Extra Hard	125° C.
Akco Resin, Extra Hard	140-145° C.
Amberol K-12-A	148-175° C.
Bakelite XR2963	138–150° C.
Beckacite 1100	127–142° C.
Beckacite 1106	127–142° C.
Lewisol N2	130–135° C.
Robert Rauh N2	135–145° C.
Q. D. No. 1	135–145° C.
Q. D. K.	140-150° C.

Hardening Rosin

Five hundred kilograms of rosin are melted in a kettle. Thirty-eight to 40 kg. of hydrate of lime are added at a temperature of 205° C., and the mixture is heated to 260° C. which causes the lime to dissolve and the mixture to clear up. The acid number of the hardened rosin amounts to half that of the colophony. In Germany the rosin is heated for some time at 175° C. Six per cent of calcium hydrate (produced from marblestone) with a magnesium oxide content of not more than 3% is then added. It is advisable to grind the calcium hydrate to a paste with a little linseed oil.

The English process, which is usually carried out in enamelled kettles consists of stirring 6% of calcium hydrate (marblestone material) into the rosin heated to 60-80° C., and it is claimed to be

possible by energetic stirring and careful operation to raise the lime additions to as high as 10%. According to another American process, 100 kg. of colophony are heated to 232° C. Six per cent of calcium hydrate is then gradually stirred into the melt within about 15 minutes and the mixture heated to 268° C. within another 15 minutes. The opinions regarding the most efficient process are thus very different. It is important to determine the most suitable percentages of lime hydrate to be added, since working by "feel" may easily cause the production of turbid material. A rosin of an acid number 145 requires the addition of 9.8% of hydrate of lime or 10.5% of zinc white. A rosin of an acid number 180 requires 11.9% of hydrate of lime or 13% of zinc white. However, the rosin must always be heated to 175° C. before adding the lime. Hydrate of lime as well as zinc white must be absolutely dry. The lime hydrate should be freshly slaked, free from carbonic acid and finely dispersed, and it is always advisable to grind this material with a little linseed oil. Most rosins require only 6% of hydrate of lime or of zinc white (Green seal) free of carbonate. It is also possible to add both materials at the same time, as for instance. 2% of zinc white and 4% of hydrate of lime. The zinc white is added at a temperature of 220 to 240° C., the mixture boiled clear, the hydrate of lime added and the mixture heated for some time at 275° C. If the hydrate of lime contains more than 3% of magnesium oxide, the melt thickens.

According to the Haines Process, the lime rosin can be boiled directly with oil, satisfactory results being obtained with two different methods of application: The oil is either boiled with the whole of the rosin at once or only with part of it, the remainder being added later in form of a lime rosin-benzene solution 1:1. The results of this process are as follows: The viscosity of the pigmented varnishes decreases if larger quantities of the rosin are boiled directly with the oil. Skinning of the pigmented varnishes decreases in the same The larger the quantities of manner. rosin directly boiled with oil the more pronounced is the whitening of the varnishes in touch with boiling water and the slower is the disappearance of the whitening. The behavior of the products towards cold water is similar. Maximum adhesion after 24 hours of storing in water is exhibited by a varnish, half of the lime rosin contents of which had been boiled with oil. This varnish also exhibits the largest pressure resistance; it is also superior in its behavior towards rapid weathering while if subjected to normal weathering conditions, the gloss of the clear varnish decreases directly with the increase of the amount of lime rosin boiled directly with oil.

The larger the quantity of lime rosin boiled with oil, the more pronounced becomes the sensibility of the product towards subsequent covering of the film

with nitrocellulose lacquers.

The Koehler process for the direct boiling of lime rosin with oil is carried out as follows: The necessary quantities of rosin are disintegrated and dissolved at 80 to 100° C. in benzene (crystal oil). At a temperature of 105 to 110° C., 4 to 5% of hydrate of lime, free of carbonic acid and lumps, and suspended in benzene, is added. The kettle, or boiler, must not be filled to more than onefourth of its capacity since the process is accompanied by strong foaming of the contents. The temperatures must not rise above 120 to 125° at the most. After foaming has subsided, the varnish can be produced at once by adding acid-poor, water-clear stand oil (with an acid number of not more than 20), tapping the mixture and centrifuging. If white enamels are produced, lime rosins must be employed which are made from excelsior rosin. One day after the production of the varnish 17 kg. of zinc white and 18 kg. of lithopone are added per 60 kg. of varnish. The mixture is thoroughly stirred and left to stand for at least 2 days. Five kilograms of varnish and 2-3% of cobalt siccative are then added and the product thinned in accordance with requirements. The subsequent addition of varnish tends to improve the gloss of the product. One to 2% of gloss-improving substances may also be added if necessary. If top grade enamel

varnishes are to be produced it is advisable not to add linseed oil-stand oil alone, but also about 20% off wood oil-stand oil. However, both types of stand oil are to be boiled separately since if the two oils are boiled in common, the wood oil would thicken before the linseed oil had been boiled sufficiently.

Investigations towards improving the hardening process have led to the following formulae: 100 kg. of colophony are heated with 1 kg. of cadmium oxide to 200-250° C., stirring continuously. After complete solution, 5 kg. of hydrate of lime are added and the product left to cool down to room temperature. A very satisfactory lime rosin varnish is

obtained by this process.

Another process is the following: 0.5 cc. of 33% caustic soda solution and 45 g. of paraformaldehyde are added to 100 g. of crude cresol heated to 80 to 100° C. As soon as the paraformaldehyde has been dissolved, the mixture is cooled and added to 800 g. of colophony heated to 200-250° C. The mixture is then stirred until the smell of phenol has disappeared. One hundred grams of this alcohol-soluble product is treated with 1 g. of precipitated or fused lithium resinate, the product obtained being easily soluble and free of separations.

A number of important guiding rules have to be observed in the production of glycerin-rosin esters. Esterification is almost universally effected in apparatus with reflux coolers, the operating temperature being about 250° C. The amount of glycerin added exceeds by about 3% that determined by calculation from the acid number of the resin. Esterification is complete after about 3 to 5 hours. The temperature is then increased to 300-320° C. in order to drive off the excess glycerin, the water of reaction and the volatile constituents of the resin. It is recommended to add 0.5% of boric acid which accelerates the esterification and prevents re-saponification by the water of reaction.

Investigations carried through in the State Industrial Research Laboratory at Tokyo (Japan) resulted in the following discoveries: (1) If aluminum kettles are employed, this metal appears to exert a catalytic influence on the process of esterification. (2) The acid number of the resulting rosin esters drops rapidly if operations are carried on at a temperature of 200° C. (3) Fifteen to 19% of the rosin is the most suitable glycerin contents. Higher glycerin contents tends to soften the product. (4) Excessively long heating causes darken-

ing of the product. (5) Dehydrating agents increase the speed of esterification. Suitable dehydrating agents are the hydrates, oxides and carbonates as well as the organic salts of metals, for instance, the formates of calcium and barium. Undesirable additions are boric acid and manganese borate. (6) A metallic salt addition raises the rosin ester

softening temperature. Typical and characteristic variation of the esterification process can also be observed in the various countries. America, glycerin-rosin esters with acid numbers up to 3 are produced in aluminum kettles. "WW-rosin" is used for light colored products. After charging the kettle it is hermetically closed and the contents melted either in a vacuum or by passing through carbonic acid. Ten to 18% of glycerin (calculated on the amount of resin used) is then added and the mixture heated for some time to 205° C. and finally to 288° C. The water vapors are permitted to escape through a reflux cooler, the glycerin flowing back into the kettle. If rosin ester of an acid number of 5 to 10 is employed, about 12% of glycerin is added. It has also been found here that an excess of glycerin tends to soften the product while excessively long heating darkens the product. The varnishes produced from glycerin ester exhibit a high gloss. They are neutral in character and resistant toward basic pigments. They do not tend to crystallize, they are free of water, flow well, but are not easily mixed with drying substances, while colophony absorb them with ease. Balm and wood colophony, as well as mixtures of the two types of colophony can be esterified. The products of wood colophony are somewhat cheaper and exhibit a lower melting point. Instead of glycerin, other hydroxyl compounds, such as naphthol or benzyl alcohol, may be used, while fossil resins can be employed instead of colo-

If, during esterizing, up to 10% of previously melted Congo or Manila copals are added, the melting points are considerably raised and the color darkened.

phony.

In Russia, rosin esters of an acid number of 4 to 5 are produced by means of catalysts, such as zinc. Rosin and zinc catalyst are jointly heated to 275 to 280° C. Eighteen per cent of glycerin is then added, the product having an acid number of 4. If catalysts are not added, it is possible by adding 24% of glycerin to obtain a product with an acid number of 25.5. A. Kogan recommends zinc chlo-

ride as zinc catalyst, while another suitable catalyst is iron trichloride in connection with hydrochloric acid gas. The original saponification number of about 173.3 is lowered by the catalytic process to about 30-40. The rosin is not appreciably changed by the use of catalysts

ciably changed by the use of catalysts. According to U. S. Patent 1,771,044, it is possible even to produce rosin esters of an acid number 1 by esterizing the rosin or the resinous acid with dichlorhydrine or dibromhydrine in presence of alkalies. For instance, 75 parts of WW-colophony are dissolved in 100 parts of alcohol containing 10 parts of caustic soda. This solution is heated to 80° C. (reflux cooling) and gradually treated with 25 parts of dichlorhydrine of a boiling point of 174° C. The mixture is then boiled 15 hours (reflux cooling), the sodium chloride produced is separated and the dichlorhydrine excess distilled off. The yield consists of 70 parts of 74° C. and the acid number 1.

An interesting French process provides for the use of wax alcohols. Natural colophony brands or resinates, hardened colophony or synthetic resins are made to react with wax alcohols, such as cetyl alcohol or cholesterol. For instance, 85 parts of colophony, 15 parts of lanolin and 2 parts of hydrate of lime are processed together. After heating the mixture of the first two constituents to 200° C., the hydrate of lime is added in small portions, and under continuous stirring, and this temperature maintained for some time. The product of reaction is transparent; it is soluble in the common solvents and yields varnish films of considerable plasticity and resistance. Another variation of this process provides for the heating of a mixture of 88 parts of colophony and 12 parts of cetyl alcohol or cholesterol to 200° C. with, or without, catalysts. Conditions are improved by operating under pressure or in an inert gaseous atmosphere. Or 85 parts of colophony are heated with 8 to 10 parts of glycerin and 5 to 7 parts of purified lanolin.

Esterization can be combined with the rosin production in the case of ester rosins as well as in operating with lime rosins. For instance, colophony is melted at 193° C., 24% of wood oil added and the temperature raised to 250° C. Ten per cent of glycerin is then added and the temperatures maintained at 288° C. for 6 hours. The kettle is finally removed from the fire and the glycerin rests removed by the addition of 5% of boric acid which forms volatile com-

pounds with the glycerin. The product, thinned with lacquer benzene, represents

a satisfactory varnish.

Glycerin rosin ester-wood oil varnishes must be boiled and cooled down rapidly. Cooling can be effected by neans of cold water or by adding cold varnish or cold linseed oil refined with alkali. Boiling with 2.5% of litharge requires the consideration of the following factors: The varnish is water resistant and impervious to gases only if boiled at 296-302° C.; the fatter the wood oil varnish, the more durable is the film, but the more pronounced is the danger of gelatinization during boiling, which can, however, be reduced by adding colophony; the lower the degree of acidity of the ester rosin or the wood oil, the greater is the danger of gelatinization and the more difficult is the addition of drying substances; the larger the ester rosin contents, the brighter and harder is the film and the more rapid is the rate of drying. Attention is called to the fact that the addition of colophony has a slightly deteriorating influence on the quality of the varnish. Addition of lin-seed oil, fish oil, soya bean oil, etc., lowers the water resistance of the film and reduces the speed of initial drying, but improves the gloss, the life and the elasticity of the film. Ester rosin varnishes must never be mixed with cold oil varnishes, as the components of this mixture do not combine with each other in the cold.

Increasing the Melting Point of Rosin

The melting point of rosin can be raised from 61 to 91° C. by 2 hours blowing with air in the presence of 1% cobalt oxide when molten, and to 107° C. after 6 hours. The amount of petroleum etherinsoluble substances (hydroxy acids) increased from 17.89 to 46.07%, the acid number and saponification number decreased from 159.56 to 145.26, and 170.00 to 161.29 respectively, and the esterification number increased from 10.44 to 16.03.

Purification of Rosin

The rosin is crushed, melted in a kettle, allowed to stand for 30-60 minutes, decanted from the impurities into a second kettle, boiled 1 hour with 20% of a 9° Bé. sodium chloride solution; the supernatant sodium chloride is siphoned off, and the treatment with sodium chloride repeated till a sufficiently light colored rosin is obtained. Soaps made from such

rosin are lighter in color than those made from unpurified rosin.

Synthetic Resin Emulsion U. S. Patent 1,999,715

One hundred parts, by weight, of ground resorcinol are placed in a metal container or kettle which is jacketed with an outside container, in turn equipped for steam heating and water cooling. permits of temperature regulation during the chemical reaction as it is very essential to carefully control the temperature of the reacting substances in order to prevent their conversion to the insoluble stage. The kettle may be equipped with suitable agitators to permit the rapid churning of the kettle contents. To the resorcinol, 112 parts, by weight, of 37.5% formaldehyde solution (formalin) are added, and the temperature is increased to a maximum of 60° C., so that the resorcinol will dissolve.

In a separate container, 8 parts, by weight, of para-nitraniline are dissolved in 20 parts, by weight, of cresol having a boiling point range of from about 215 to 230° C. The melted para-nitraniline is added to the formaldehyde solution, and the mass is thoroughly agitated, the reaction temperature being raised to from 70 to about 75° C. A plasticizer of vegetable or animal oils and wax with a filler and suitable coloring material may be added as a paste to the mixture in the kettle. As an example, 8 parts, by weight, of clay, 0.8 part, by weight, of beeswax, and 1 part, by weight, of iron oxide, all best ground in a paint mill or ball mill to obtain thorough dispersion. The paste has been found to mix readily with the thickening liquid in the kettle.

For best results, the temperature should be maintained at no higher than 80° C. When the mass in the kettle becomes stringy, and before gelatinization can take place, the emulsification enhancing agent is added. A water solution (65 parts, by weight) containing 0.1% of borax is added, first slowly, and then rapidly to the agitated resin. The borax, or any other suitable alkaline salt, serves to so enhance the emulsifying action of the wax and/or the oils (and gums, if any are used) as to enable them to properly maintain the resin in suspension in the water. This results from the fact that the borax or other alkaline salt used reduces the surface tension of the water from its normal value at the operating temperatures, thereby more readily effecting a wetting of the resin particles and enabling the wax or other fatty material used to more easily retain the resin particles in suspension. The temperature is dropped to about 20° C. and thickening of the resin takes place. A solution of alcohol (about 65 parts, by weight) or an equal quantity of a 20% benzol or toluene solution in alcohol may be added to the emulsifier to obtain proper consistency. The varnish is immediately strained to remove any foreign particles. The mass is then cooled, with accompanying increase in viscosity of the varnish, and additional alcohol or solvent mixture may be added to obtain the desired viscosity.

Flexible Synthetic Resin U. S. Patent 1,999,097

Diethylene Glycol 103 oz. Phthalic Anhydride 148 oz.

These ingredients are mixed together and heated gently in a suitable receptacle until all of the phthalic anhydride has melted, the temperature of the mix is then gradually raised to approximately 165° C, and maintained at this temperature for approximately 4 hours. The resulting resin on cooling is a viscous liquid having a light amber color and is soluble in acetone, alcohol, chloroform, nitrocellulose and cellulose acetate solution.

Synthetic Molding Resin U. S. Patent 2,010,225

A mixture of 9 lb. of asbestos, 3 lb. of shellac and 2 oz. sulphanilic acid is repeatedly passed and repassed between hot rolls maintained at a temperature sufficient to keep the shellac in the composition molten. When the mixture in this manner has been rendered uniform in distribution, the resulting plastic mass may be pressed into slabs, or so-called biscuits of any desired type or shape. These biscuits may then, if desired, be subjected to a heat curing or baking process. The exact details and conditions of any such intermediate step will, to a large extent, depend upon the nature and service to which the later manufactured article is to be put. This product may then be placed in a mold either in biscuit (by softening on a steam table) or in a powdered form, and subjected to required heat and pressure necessary for forming a hard, resistant, less fusible Where a limited amount of object. agent and previous heat treatment of the biscuit material has been employed, it l

may be necessary to cool the mold during the pressing operation. The molding may take place in a number of ways but good results may be obtained by softening the biscuit material at 300° F., placing a slight excess in the mold and subjecting the same to 2700 lb. per sq. in. The pressure is not released until the material has cooled to a temperature sufficiently low to be readily handled without deformation.

Synthetic Resin Paper Size Emulsion U. S. Patent 2,022,004

Resin A

Glycerol 15.6 oz. Phthalic Anhydride 20.18 oz. Stearic Acid 64.22 oz.

The ingredients are heated together with stirring in a suitable vessel, the temperature being carried to 200° C. over a period of 1 hour, then maintained at this point until an acid number of 47 has been reached. This requires approx-

imately 21/2 hours.

The preferred method of converting the resin into an aqueous emulsion, defined here as a dispersion of very fine particles of the resin in water, is as follows: 100 parts of Resin A at 100° C. and 61.0 parts of a 5% solution of sodium hydroxide at 60° C. are added simultaneously and in proportionate rates to 349 parts of water at 60° C., with rapid agitation during the mixing operation. The alkali solution should be added slightly in advance of the resin, and the emulsion should be stirred for a few minutes after the mixing operation has been completed. This gives a 20% emulsion of the resin. The amount of sodium hydroxide used in preparing the emulsion is insufficient to neutralize completely the titratable acid in the resin. The resin is therefore not present in the water in complete solution, but as an emulsion, i.e., it is largely in the form of a physical dispersion in the water. This is a very substantial difference from those cases in which the resin is completely neutralized, as in the prior art. The suitability of the present emulsions enables one to use resins which are carried to a lower acid number and, hence, a more complete resinification. Lower acid numbers and higher resinification are necessary to give the improved water resistance when applied for the purposes of this invention. High acid number resins require alum in addition to alkali to develop their maximum water resistance; the use of more completely esterified products obviates this disadvantage. The emulsion can be diluted with warm water to any desired concentration.

Plaster of Paris Synthetic Resin Casts British Patent 425,742

Plaster of Paris casts are impregnated with an aqueous solution of the reacting components of phenol formaldehyde resins in the early or molecular stages of condensation to increase their hardness. toughness and gloss, and to secure their impermeability to water. The product is capable of taking a high polish, and of being stained. Instead of phenol, cresol and homologues thereof may be used. In an example a plaster cast is immersed until saturated in a mixture of equal weights of commercial cresol and 40% formaldehyde and 1 or 2 parts of a 50% solution of potassium hydroxide. solution is warmed to 35° C. The object is then stoved at 100° C.

Synthetic Dielectric Resin Canadian Patent 342,586

Abietic acid 800, glycerol 770, phthalic anhydride 852, ethylene glycol 965 and linseed oil acids 80 parts by weight are heated under reflux to 175-180° C. for approximately 30 minutes, and 320 parts by weight of tung oil is added in 4 parts. Succinic acid (1820 parts) is added and the mass cooked until a resin is formed. The excess of glycerol is removed by vacuum distillation. The resin is used in coating compositions for fibrous material as cloth and paper in order to impart a flexible, tough film of good dielectric value, unaffected by mineral oil or petroleum or aromatic solvents.

"Albertol" Type Synthetic Resin Formaldehyde 0.85 l. Phenol 1 kg. Hydrochloric Acid 0.02 kg. Reflux 2 to 3 hours.

Pour off liquid and dry residue in

vacuo at 100° C.

To 0.3 kg. of above resin add 0.7 kg. rosin and heat to 120-130° C. When solution is complete add 0.4% calcium oxide and heat to 290° C. Maintain at this temperature until a sample is soluble in oil and has an acid number of about 30.

"Haveg" or "Prodorite" Type Materials

An acid proof material suitable for tanks and other apparatus is made of

80% sand, an appropriate amount of coal or oil bitumen and of 5% acid resistant minerals (grog, clay, etc.); the mixture is heated to 150-200° F. and molded to the desired shape. It sticks to iron, is resistant to hydrochloric acid and to diluted nitric acid. Coumarone tar can be used as a protecting varnish for low temperature and for molded objects of a low mechanical strength. "Haveg" from asbestos and bauxite has a mechanical strength similar to that of cast iron.

Sound Record Composition British Patent 408,969

A particularly suitable resin is formed by the conjoint polymerization of vinyl chloride 80 and vinyl acetate 20%. The resin may be mixed with a filler, e.g., wood filler, cotton flock, silica, mica or with a plasticizer, e.g., dibutyl phthalate, tricresyl derivative, glycol, glycerol esters

Gramophone Record Composition

Lac	15	oz.
Copal	1.5	oz.
Silica	19	02.
Barytes	19	oz.
Carbon Black	5.5	oz.
Scrap	40	oz.

For cheapness, part of the carbon black is often replaced by mineral black.

The scrap is spew and rejected records, etc. The amount of lac varies, dependent upon the grade used, it being generally considered that T.N. Orange is about the lowest that can conveniently be employed at present.

Vinyl Resin Canadian Patent 352,766

Polymerize following a	ıt	about 40	°C	١.
Vinyl Chloride		80	oz.	
Vinyl Acetate		20	oz.	
Hexane		100	oz.	
Benzovl Peroxide		0.5	oz.	

Vinyl Acetate Resin German Patent 615.995

Water	200 g.
Vinyl Acetate	200 g.
Hydrogen Peroxide (30%)	
Soda Ash	1 g.
Heat at boiling point for 1	to 2 hours.

T۸

Bleaching Beewax

7.0		
<i>a</i>	Water Potassium Bichromate	70–75 cc. 15 g.
u.	Water Potassium Bichromate Sulphuric Acid (60° Bé.) Boil.	15–20 g.

add

b. Beeswax, Molten 100 g.

Stir until color becomes greenish blue. Cool. Remove solution shortly before wax solidifies. Boil wax with clean water to remove acid.

Synthetic Beeswax

U. S. Patent 1,983,672

Formula No. 1

Five hundred grams of a mixture of the higher paraffin hydrocarbons melting at 74-76° C. (Superla wax) is mixed with 10 g, of manganese cleate and oxidized in a glass reaction vessel at 130-140° C. by oxygen passed through the hydrocarbons by means of a tube with many small orifices submerged in the hydrocarbons. The oxygen is passed through the hydrocarbons at the rate of approximately 1/2 cu. ft. per hour. At the end of 144 hours the contents of the vessel has gained in weight about 20 g. It has an acid value of about 23 and an ester value of approximately 100. In physical properties this product closely resembles beeswax except it melts at a temperature approximately 10 degrees above the melting point of true beeswax.

No. 2

Two batches of 1500 g. each of the ozokerite wax ("Utahwax") with a melting point of 73° C. are mixed with but 1% of their weight of manganese oleate and then oxidized simultaneously in 2 flasks A and B. Dry oxygen at the rate of 3/10 cu. ft. per hour is passed into the flask A and brought into intimate contact with the hydrocarbon therein. The oxygen and the vapors coming off from the first flask A are passed through a soda-lime tower and then into flask B. The temperature of each flask is maintained at approximately 120° C., and after oxidation for 288 hours the reaction is discontinued. The product in each flask resembles commercial beeswax and is suitable for use as a beeswax substitute. The acid value of the product in flask A is about 25.8 and its ester value about 50.6. The product in flask B has an acid value of about 46.7 and an ester value of about 56.6.

Raising Melting Point of Montan Wax U. S. Patent 1,966,168

Formula No. 1

Crude montan wax with a melting point of 80° C. is fused. Two-tenths per cent of calcium hydroxide suspension is added to fused wax while continuously stirring, the temperature being slowly raised up to 90°. Stirring is continued at this temperature for about half an hour. In this way the melting point of the montan wax is raised to 85°.

No. 2

Crude montan wax having a melting point of 80° is fused and 0.2% of calcium hydroxide is introduced at a temperature above 100° while continuously stirring until uniform distribution has taken place. After about half an hour treatment the melting point of the wax is raised to 85°.

No. 3

Crude montan wax solution obtained in the course of manufacture is mixed with 0.2% of calcium hydroxide, care being taken that uniform distribution takes place. After the hydroxide has acted for about half an hour the melting point of the wax raised about 5°.

"Hardened" Stearic Acid Wax

Stearic Acid 75 oz. Magnesium Oxide 5.3 oz.

Heat with stirring for ½ hour at 130-150° C. Pour at lowest possible temperature.

Illumination Candles

Paraffin (50-52° C.)	79 g.
Stearin	19.5 g.
Carnauba Wax, Bleached	d 1.5 g.

Wax Lighting Tapers

Paraffin Wax (40-42°	C. or	
42–44° C.	65-85 g.	
Ceresin (58-60° C.)	30-10 g.	
Beeswax	2-3 g.	
Turpentine, Thickened	3-2 g	
TTT: 1 . C 1	Alaman Ja 20	,

Wick of loose cotton threads, 30 together for a size of 2-4 mm., wound on wire.

Lor	ng	Burning	Candles
U.	s.	Patent	1,954,659

Paraffin Wax 49 lb. Hydrogenated Vegetable Oil 51 lb.

Molded Candle

U. S. Patent 1,960,994

Beeswax	70	oz.
Stearic Acid	20	oz.
Paraffin Wax	10	oz.
"Cellosolve"	1	٥z.

Sealing Wax for Candle Decorations Rosin 50 g. Ruby Shellac 3 g. Gypsum 1 g.

Dental Wax

Stearic Acid	1 lb.
Paraffin Scale Wax	2 lb.
Glyceryl Tristearate	1 lb.
Carnauba Wax	2 lb.
Ethylene Glycol Glyceryl	
Stearate	2 lb.

Ceresin Wax

Ceresin wax consists of a mixture of ozokerite and paraffin waxes.

Starting with pure yellow ozokerite and melting together in the following proportions with paraffin wax gives the following blends:

Pure Ozo-

I UIC OLO		
kerite Wax	Paraffin	
White	Wax	
M. P.	M. P.	gives
75° C.	50° C.	Ceresin Wax
4 oz.	1 oz.	M.P. 73.5° C.
4 oz.	2 oz.	M.P. 71.7° C.
4 oz.	3 02.	M.P. 72.5° C.
4 oz.	4 oz.	M.P. 69.7° C.

When pure white ozokerite is used the following results:

Pure Ozo-		
kerite Wax,	Paraffin	
White	Wax	
M.P.	M.P.	gives
75.7° C.	58.3° C.	Ceresin Wax
4 oz.	1 oz.	M.P. 74.4° C.
4 oz.	2 oz.	M.P. 73.2° C.
4 oz.	3 oz.	M.P. 72.5° C.
4 oz.	4 oz.	M.P. 72.0° C.

Electrotypers' Waxes Formula No. 1

Beeswax	$5\frac{1}{2}$	lb.
Paraffin Wax	3	lb.

Burgundy Pitch	34 lb.
Rosin W.W.	½ lb.
Zinc Oxide	1½ lb.

Melt together the waxes and resins and add the zinc oxide slowly with good mixing.

n	ixing.	011.0	0 510 1123	******	8,,
		No.	2		
	Ozokerite	±10		631/2	lh.
	Beeswax			$31\frac{3}{4}$	
	Graphite Powd	er		$4\frac{3}{4}$	
		No.	3		
	Beeswax			85	lb.
	Burgundy Pitcl	1			lb.
	Turpentine	-			lb.
	•	No.	1	7.7	
	Ozokerite	110.	*	05	lb.
	Graphite Powd	ar			lb.
	Grapino I owa	C.		J	10.
		No.	5		
	Ozokerite, Gree	n		33	lb.
	Paraffin Wax			50	lb.
	Rosin W.W.			16	lb.
	Petrolatum			1/3	lb.
		No.	6		
	Ozokerite, Brov	vn		.31	lb.
	Graphite Powd			2	lb.
	Pine Pitcl.				lb.
	Rosin Oil			1/4	lb.

Insulating Wax

Carnauba Wax	1	lb.	14	OZ.
Yellow Beeswax			4	OZ.
Venice Turpentine			6	OZ.
Gum Obsidian			-	OZ.
Sulphur	2	lb.	8	OZ.

Cook until thoroughly uniform. This wax should have a melting point of 285° F, and a flash point of 499° F.

Recording (Phonograph) Wax

Formula No. 1

Stearic Acid			84	1b.
Melt and add	slowly	with	stirrin	
Litharge			81/2	lb.

Boil off water at 220-230° F. Stirring must be of such type to prevent caking at bottom of kettle. When solution is complete add slowly (by sifting in):

Soda Ash 7 lb.

When a drop cools to a clear mass reaction is complete. Drive off all gas, froth and water by heating up to 270° F.

If a brown wax is desired add to above

Stearin Pitch 2 lb.

If a black wax is wanted add some oilsoluble nigrosine to brown formula.

No. 2		
Litharge Soda Ash Paraffin Wax	4½ 4 30	lb. lb.
Follow method exactly as in No. 1.	ı Fo	rmula
No. 1.		
Shoemakers' Sewing W Candelilla Wax Rosin	2	Ib. Ib.
Burgundy Pitch		lb.
Rosin Oil		ib.
Lard		lb.
Mineral Oil (Heavy)	1	lb.
Shoe Finishers' Black Stiel Candelilla Wax Rosin Carnauba Wax (North Country) Oil-Soluble Black Dye Carbon Black Paraffin Wax	9 1 32 6 1/4	lb. lb. lb. lb. lb.
Black Padding Wax		
Carnauba Wax (North Country) Ozokerite (Green) Paraffin Wax Rosin Oil-Soluble Black Dye	2 58 2	lb. lb. lb. lb. lb.
Tree Grafting Wax		
Wool Fat, Neutral Rosin Ceresin (58–60° C.) Beeswax Rosin Oil	4(

Wax for (Wounded) Trees

Formula	Ma	1	
r ormula	TAO.	1	

Rosin			60 g.
Alcohol			40 cc.
35-14	+1	0 dd 4h a	.11

Melt up the rosin, add the alcohol cau tiously. Stir until cold.

No. 2

Melt up:		
Rosin	15	٤.
Linseed Oil		čc.
Turpentine (Thick)	1	cc.
Yellow Beeswax	2	g.
Melt together below 78° C.		0
Add:		

Fill into air-tight cans.

Alcohol

Non-Inflammable Film U. S. Patent 1,981,132

4-5 cc.

	,	•		
Cellulose Acetate			100	lb.
Triphenyl Phosphate			20	lb,
Diethyl Phthalate			10	lb.

Transparent Foil or Film Base British Patent 411,471

Cellulose Acetate		
(Anhydrous)	100	lb.
Acetone (Anhydrous)	400	lb.
Diethyl Phthalate ,	16.7	lb.
Diacetin	5	lb.
Triphenyl Phosphate	8.3	lb.

Polychromatic Printing Plate U. S. Patent 1,999,549

Dextrin 10, glycerol 10, soap 10, talc 10, naphthalene 0.5 and water 16 parts are mixed with a pigment.

WAX TYPE ACIDS AND HIGHER WAX TYPE ALCOHOLS

					10,	11 112	1110, 1	LIA	DIICO					212	2
Occurrence	Free in beeswax, montan wax, earnauba, also as cerotate in insert wax, wool wax, and carnauba.	Free in montan wax.	Free in beeswax and montan wax.	As tri-palmitin in palm oil and Ja- pan wax; as cetyl palmitate in spermaceti; as myricyl palmitate in beeswax.	As laurin in coconut oil and Japan wax.	As myristin in coconut and palm- nut oils.	As cetyl palmitate in spermaceti.	Spermaceti.	As ceryl palmitate in opium wax, as ceryl cerate in Chinese insect wax.	As myricyl palmitate in beeswax, carnauba, sugar cane wax.	Carnauba wax.	Cochineal wax.	In wool-fat and sperm oil.	Plant cholesterol.	
Soluble in	Warm Alcohol	Methyl Alcohol	Alcohol Ether	Alcohol Ether	l	1	Alcohol Ether Benzol	I	Alcohol	Ether Alcohol	Ether	1 1	Ether	Benzol	
Specific Gravity at 15° C.	.836 at 79° C.	1	.847	.846	I	l	.810	Ĭ	I	Į	I	1	J	1 1	
Melting Point	77.8° C.	83 ° C.	91 ° C. 70.5° C.	62.2° C.	43.5° C.	53.8° C.	50 ° C.	59 ° C.	79 ° C.	88 ° C.	103 ° C.	103 ° C.	147 ° C.	137 to 138° C. 134° C.	
Formula	$\mathrm{CH}_3[\mathrm{CH}_2]_{24}\mathrm{CO.OH}$	$ m CH_{3}[CH_{2}]_{26}CO.OH$	С ₃₀ Н ₆₁ СООН СН ₃ [СН ₂] ₁₆ СООН	$\mathrm{C_{16}H_{32}O_2}$	$ m C_{12}H_{24}O_{2}$	$\mathrm{C_{14}H_{28}O_{2}}$	$\mathrm{C_{16}H_{33}OH}$	$C_{18}H_{37}OH$	$C_{26}H_{53}OH \\ C_{27}H_{55}OH$	$ m C_{30}H_{62}(OH)_2$	$C_{24}H_{48}(\mathrm{OH})_2$	$\mathrm{C}_{30}\mathrm{H}_{60}(\mathrm{OH})_2$	$\mathrm{C}_{27}\mathrm{H}_{44}\mathrm{OH}$	-	
Waxy Material	Cerotic Acid	Montanic Acid	Melissic Acid	Palmitic Acid	Lauric Acid	Myristic Acid	Cetyl Alcohol	Octodecyl Alcohol	Ceryl Alcohol	Myricyl Alcohol	Anonymous Alcohol	Cholesterol or Cholestery Al.	cohol	Iso-Cholesterol (Isomeric) Phytosterol	

WAXES
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L PROPERTIES (
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L AND CHEMIC
AND
PHYSICAL

Нудгосягропя	2-56% 5-75% 4-55% 9-50% a 50% a 81.3% a 80%	2%	,		13% 14%	1
Alcohols and	以 品 型 全 D D	ස් 	1	11	39–43% 33–44%	1
Acety Val		•	l		n 1 2	63
Ratio Acetyl No. Val.	.68 3.6–3.8 5.7 4.7 or 39 31 29.3 ————————————————————————————————————	0	0	11-35		
Fatty Acid %	47.8 33.3 29.0 48.0 51.5 — — 11-15 56-64 (Dist.)	0	0	06	53.5	
XES Aeid est Val.	20 12 10-20 2.5 3 3 - 54.5 15-20 15-20 73-83	0	0	6-20	1.5	12.2 50%
N WAJ Sromine srmal Te	.26 1.3-2.0 1.7-2.4 02	1	ı	.6-2.6	14.3–15 — — .5–30	3-4.6
IE COMMON WAXES Lodine Bromine Acid % Val. Thermal Test Val.	2.0-4.0 88 85 35 12.5-15.0 0-1.5 11-13 9-17 10-15 16-20	0	0	5.0-16.0	86–91 — 3.0–4.0	25-43
TIES OF THE Sap. Unsap. 1 No. Matter %	55.5 69.0 74.0 55 49.5	100	100	.7–15	40 — — 51.5	
ERTIES g Sap.	206-216 90-101 80-90 50-65 67-88 82-93 1150-160 30-45	0-1.3	1.3	219–237	66–76 88–93 77–79 122–134	101-104
PROPEI Setting Point	60-63 63-68 80-87 80-81 80-81 70-80	*	92	1 1	1, 1 1 1	I ,
EMICAL M.P. °C.	41 62-66 58 66-70 83-84 81-83.5 62-70 73-84	96-56 35-75	92-69	52–59 82.0	$\frac{12}{9}$	37-41
PHYSICAL AND CHEMICAL PROPERTIES OF THE COMMON WAXES Sp. Gr. Ref. Index M.P. °C. Point No. Matter % Val. Thermal Test V	1.440–75° 1.456–75° 1.472–43°	.870910 1.4331-1.4450 26-56	.913923 1.4415-1.4464	1.4518	1.459 1.462–25° 1.456–25° 1.4198	1.480
PHYSICA Sp. Gr.	.993997 .960947 .980 .982972 .982970	.870910	.913923	.976993	.879 .876 .878 .932–.963	.945
	Bayberry (Myrtlewex) Not a true wax Beeswax Cane Sugar Wax Candelilla Wax Carnauba Wax Chinese Insect Wax Cotton Seed Wax Flax Seed Wax Montan Wax	Paraffin Wax Not a true wax	Ozokerite Ceresin Wax	Japan Wax Not a true wax Raphia Palm Wax	Sperm Whale Oil: Head Oil Body Oil Arctic Sperm Oil Spermaceti	Wool Wax

					RU	BBER,	RE	SIN	s, w	A.	XF	S, PLA	STICS	3		321
	1	1			Solubility in Fusel Oil	Soluble	1	Soluble		-	1	Soluble	1	ı	Soluble	Soluble
	- 25-200	.1-0.2	9.5 .01		Solubility in Carbon Tet- rachloride	Soluble	1	Soluble			١	Soluble	1	1	Soluble	Soluble
	Variable 95	130-186 —	.98 96–99.5		Solubility in Turpentine	Soluble	İ	Soluble			1	Soluble	1	ı	Soluble	Soluble
	10	55-180	.5-30	AXES	Solubility in Petroleum Ether	Insoluble	1	Cold—insolu-	Hot-soluble		1	Soluble	1	1	Soluble	Soluble
OF WAXES	198 .5-2.0	147-180 5-15	200 0.5	SOLUBILITY DATA OF COMMON WAXES	Solubility in Ether	Soluble in hot I and cold	ı	Cold—insolu- C	-soluble		I	Soluble in cold and hot	, 1 ,	1	Soluble in hot and cold	Soluble in hot and cold
ADULTERANTS OF WAXES	30-60	over 100 — 1	49–56 —	ITY DATA 01	Solubility in Chloroform	Cold—insolu- ble Hot—soluble	1	Cold—insolu-	Hot—soluble		1	Soluble in cold and hot	1		Soluble in sold and hot	Soluble in sold and hot
A	36	- 000	1.4380 46	SOLUBIL	Solability in Acetone	Insoluble in cold Soluble in hot	1	Insoluble in	Not very sol- uble in hot		1	Insoluble in lot and cold	*		Insoluble in cold and slightly soluble in hot	Insoluble in cold and sol- uble in hot
	Oil —	1.07–1.08	1		Solubility in Hot Acetic Anhydride	Melts, floats, dissolves—so-lidifies on cooling	o	Becomes			1	Dissolves and solidifies on cooling		Dissolves and solidifies on cooling	1	j.,
					Solubility in Alcohol	76° C.	63° C.	82° C.			Insoluble	76° C.	Dist. Montan Wax 70° C.		Insoluble	44° C.
	Hardened (Hydrogenated)	Rosin	Stearin			Beeswax	Candellila	Carnauba		Chinese Insect	Wax		Montan	Ozokerite	Paraffin	Spermaceti

SOAPS, CI	LEANERS
Solvent Liquid Soaps Formula No. 1	No. 9 Soap
1:1 Hexalin-Methyl Hexalin 250 kg. Potash Lye (50° Bé.) 270 kg. Water 1300-1800 kg. The ingredients are stirred together in	Coconut Oil 21 — 6 kg. Soya Bean Oil — 8 12 kg. Potassium Hydroxide Solution (50%) 9.5 4.6 9.6 kg. Sugar 12 8 — kg.
Hexalin or methyl hexalin may be partially replaced by other solvents as shown below: No. 4 Linseed Oil 184 kg. Hexalin 275 kg. Potash Lye (50° Bé.) 73.5 kg.	Potassium Carbonate — 2 — kg. Water 55.5 71.4 60.2 kg. Oil of Lavender — — 0.1 kg. Linalyl Acetate — — 0.1 kg. The oil is first run into a pan fitted with an open steam coil which serves to
Water	both heat and agitate the pan contents. Heat the oil to about 70° C. and gradually add the caustic potash solution until the oil is completely saponified. It will be found necessary to add water before all the alkali has been introduced.

This is one method of checking foaming Potash Lye (50° Bé.) kg. which is likely to occur particularly in 615 kg. the case of cotton-seed oil and to a lesser kg. Carbon Tetrachloride 120 extent when coconut or palm kernel oil Similarly, equal weights of benzine or is used. When saponification is complete high-boiling petroleum distillates may be add sugar, glycerin, etc., and finally adused in place of carbon tetrachloride. just the water content. Allow to cool No. 6

> 60 kg. 2 kg.

30 kg.

25 kg.

45 kg.

Water

Soap

Water

Potassium Carbonate

Trichloroethylene

No. 8

somewhat, then add color and perfume if 35 kg. Soap required. Cyclohexanol 10 kg. Where possible it is an advantage to Water 55 kg. use soft water, as salts of hard water No. 7 result in the formation of corresponding 28 kg. Soap insoluble metallic soaps, which deposit or 10 kg. Trichloroethylene give a cloudiness in solution.

Liquid Soap Shampoos

Liquid soap shampoos are best made from olive oil potash soap dissolved in hot 80% alcohol in which it is completely

soluble, although the solution becomes slightly clouded on cooling. Dissolve the soap (1 part) in alcohol (4 parts) in a vessel which can be heated on a water bath and so constructed that alcohol is not lost by volatilization. When completely dissolved add coloring matter and perfume.

The formulae given are only a very few of the many that are available. Even using the same constituents of a given formula, the number of combinations could be varied in relation to fatty acid content, etc. Obviously the relative percentages of oil and alkali required for saponification would vary only between

narrow limits.

Production of Liquid High-Content Potassium Soaps German Patent 613,224

Formula No. 1

2 0111111112 2101 2		
Olein	350	g.
Coconut Oil Fatty Acid,		
a. Coconut Oil Fatty Acid, (free from Stearic		
Acid) Distilled	50	g.
₹ Alcohol	$\frac{50}{150}$	cc.
Water	210	cc.
b. Water Potassium Acetate Caustic Potash (48° Bé.)	5 0	g.
(Caustic Potash (48° Bé.)	190	cc.

Mix the two solutions. Soap contains 40% free fatty acid, is liquid down to 0° C. and gives no jelly on standing.

No. 2

	Fatty Acid of Low-Boil- ing Fraction of Sperm (Whale) Oil Cocoanut Oil Fatty Acid (Low Titre) Distilled		
a.	(Whale) Oil Cocoanut Oil Fatty Acid	1000	g.
	(Low Titre) Distilled	220	g.
ъ.	Adipic Acid Alcohol	$\begin{array}{c} 75 \\ 450 \end{array}$	cc.
c	Caustic Potash (48° Bé.) Water	$\begin{array}{c} 630 \\ 700 \end{array}$	ec. ec.

Mix α and b, and add c, with stirring. Clear, liquid soap with 40% free fatty acid.

Liquid Soap (15%)

Coconut Oil	12	kg.
Castor Oil	4	kg.
Potassium Hydroxide		
(50° Bé.)	8	kg.
Water	76	kg.
Potassium Chloride	0.5	kg.

Saponify with warming; allow to stand for 1-2 weeks, separate clear liquid by siphon, filter sludge through a Seitz filter, put both together; optionally use alcohol or glycerin.

Liquid Olive Oil Soap

Two hundred and twenty-seven kilograms of potash are dissolved in the minimum quantity of water, and into the solution is stirred a mixture of 182 kg. olive oil, 362 kg. palm oil and as much coconut oil previously warmed to 49° C. Alcohol is next run in (170 l.) and the liquid heated to 82° C. (under reflux it is presumed). After saponification and cooling, 5.6 l. water are run into the alcoholic soap.

Liquid Coconut Oil Soap

Six kilograms potash are dissolved m 20 l. water and the solution run into 20 kg. coconut oil warmed to 49° C. After adding 2.5 l. of alcohol the mixture is kept at 82° C. to saponify, when it is left to cool for 24 hours. Eighty liters of water are then added, with a little sugar, potassium chloride or glycerin if necessary.

Glycerin Liquid Soap

Thirty-five parts of good soft soap are well mixed with 21 parts glycerin, and 7 parts of water well crutched in. This is followed by 14 parts alcohol. This solution is subjected to a fairly long sedimentation after adding tale or pumice. If excessively alkaline it must be first corrected by the addition of oleic acid. Perfuming or coloring can be done if desired.

Liquid Soaps

1		
Coconut Oil	10	kg.
Castor Oil	5	kg.
Lard Oil	2	kg.
Caustic Potash (31/2 parts		0
solid)	$16\frac{1}{2}$	kø.
Water		suit

This should be easy to make. Warm up the mixed oils and add the caustic solution. Heat gently. When clear and bright, like syrup, add sufficient distilled water to the consistency required, using phenolphthalein solution (½%) to correct.

Another mixing that will not lather as readily as the previous one, but which has the advantage of being an excellent cleanser, the power of which is only slightly diminished even in hard water, is as follows:

Lard Oil, Olein or Castor Oil 50 kg.
Glycerin 150 kg.
Caustic Potash Solution
(38° Bé.) 20 kg.
Carbonate of Potash Dissolved
in 5 parts of Hot Water 3 kg.

This can be perfumed slightly and the following should give a delicate, yet pleasing, result:

Lavender Oil 2 kg.
Bergamot Oil 1 kg.
Geranium Oil 1 kg.
Patchouli Oil ¼ kg.

About 1% of this should be sufficient to give the desired effect. The nethod of making the above soap should follow along the lines described and should present no difficulty.

Formaldehyde Soap Solution

	Soft Soap	40 lb.
	Alcohol	30 lb.
	Formaldehyde	20 lb.
	Distilled Water to make	100 lb.
	As to perfume, oil of la	avender (about
1	lb.) may be added.	•

Liquid Disinfecting Soap

~	Coconut Oil	18	kg.
<i>a</i> .	Coconut Oil Soya Oil	2	kg.
b.	Caustic Potash (38° Bé.)	12	kg.
	Water, Soft	68	kg.
٥. ٠	Water, Soft Potassium Chloride	0.5	kg.

Mix a, saponify with b, dissolve in c. Prepare:

Turkey Red Oil (70%) 3 kg.
Phenyl-p-Hydroxy

Benzoate 20–25 dg.

The solution d is enough for 100 kg. of above made soap-base.

Add perfume.

Disinfectant Scrub Soaps

Cheap disinfectant soaps in England ordinarily consist of suitable tar acid derivatives emulsified in a solution of rosin soap. Creosote, phenols, cresols and naphthalene are the usual disinfectant agents. The following directions are for the preparation of liquid disinfectant soaps suitable for scrubbing floors, etc.:

Formula No. 1

Ground Rosin	17 lb.
Caustic Soda, 30%	3 lb.
Water	5 gal.
Crude Cresol	3 gal.

Boil the caustic soda in 1 gal. of water and add the rosin gradually to this. When dissolved and partly saponified, add 2 more gal. of water with continuous boiling and stirring. Add 2 gal. of cresol with stirring, then the remainder of the water and cresol. Keep covered until cold.

No. 2	
Water	61/2 Db.
Powdered Rosin	33/4 lb.
Powdered Soda Ash	1 lb.
Powdered Naphthalene	34. lb.
Filtered Creosote	½ lb.
Soft Soap	1/4 lb.
D: 1 11 1	/4 10.

Dissolve the soda ash in water and heat to boiling. Add the rosin and heat until saponified. Mix the soft soap and naphthalene separately and add the crossote to this. Add the mixture to the rosin soap with continuous stirring.

Pine Oil Scrubbing Soap

Corn Oil Soap	50 lb.
Pine Oil	10 lb.
Diglycol Laurate	5 lb.
Alcohol	3-5 lb.

Mix until uniform. A transparent jelly like product is formed.

Liquid Pine Oil Soap

		T. OIL	nuia No.	T		
Pine (Oil				300	kg.
Soya (Oil	Fatty	Acid		100	
Water	•	•				kg.

Warmed gently to be liquefied, then add Caustic Potash (50° Bé.) 40 kg. Clear Soap Oil, 1 part mixes with Turpentine 4 parts

Benzoline or 4 parts

Carbon Tetrachloride 4 parts

Dichloro-Ethylene

with water.

Dichloro-Ethylene 4 parts
or

Naphtha 4 parts to clear oils, which give excellent emul sions in water (1:1 to 1:2).

Above made Pine Oil Soap
Pine Oil
Spindle Oil Passad

12.5 kg.
12.5 kg.

Spindle Oil, Refined, 2° Engler, at 50° C. 75 kg. yields clear oil, gives excellent emulsions

No. 2

Melt
Rosin WW-F/G
Soya Oil Fatty Acid

Add

Pine Oil 105 kg. Take off 40 kg. and keep aside. To the remaining 110 kg. add:

Water Caustic Potash (50° Eé.) 15 kg. 15 kg.

1	SOA	PS,	(
Stir until glassy-transparent, above mentioned 40 kg.	add	th	e
To the product add			
Water less than			
(a tough, jelly like soap p	aste)		
or Pine Oil	100 1	kg.	
(water soluble, liquid soap)	_	
No. 3			
Pine Oil Jelly Soap			
Sova Oil Fatty Acid or			
Linseed Oil Fatty Acid	40 1	kg.	
Pine Oil	25 1	kg.	
Warm gently.			
Add:			
Water	15 l	kg.	
Caustic Potash (50° Bé.)	8 1	ζĊ.	
Caustic Potash (50° Bé.) Caustic Soda (36° Bé.)	12 l	kg.	
Water (optional) 15	-30 I	kg.	
No. 4			
Pine Oil, "Soluble"			
Soya Oil Fatty Acid or			
Linseed Oil Fatty Acid	25 1	kg.	
Pine Oil	35 I		
Warm gently in		-	
Water	10 1	ko.	
Caustic Potash (50° Bé.)	10		
Pine Oil	160		
		_	
Pine Oil Cleaning Past	te		
Glycol Laurate		lb.	
Pine Oil	25		
	1	•	

Pine Oil Clean	ing Paste
Glycol Laurate	5 lb.
Pine Oil	25 lb.
Mix and add to fo low	ing while stirring
0	E0 116

Soap Paste Paint Cleaner
Soap Chips 20 oz.
Mineral Spirits 10 oz.
Water 69.3 oz.
Oil of Sassafras 0.7 oz.

This is a semi-solid or heavy soap paste, white and permanent. It is very effective as a cleaner for painted surfaces. It is also used as a cleaner for carpets and rugs. The soap is allowed to soak in the water which is then heated to bring all the soap into solution. Same is then agitated vigorously while the mineral spirits is added and then the oil of sassafras.

Waterless Soap

11 2002 2002 1000	·r
Oleic Acid	4 lb.
Turpentine Substitute	1 lb.
Industrial Spirit	2 lb.
Castor Oil	1 lb.

Neutralized with a solution of caustic potash (1:1), 2 of water added to form

a paste and 15% of powdered borax incorporated.

Soap Powders Formula No. 1

Formula No. 1				
Palm Kernel Oil Fatty Acid 3	lb.			
or				
Tallow				
Hard Fat Bone Fat Acid 2				
Bone Fat { Acid 2	lb.			
Palm Oil (Bleached) J				
Caustic Soda (36° Bé.) 3	lb.			
Soda Ash 12	lb.			
No. 2				
Soft Soap Fatty Acids 6-7	1h.			
Hard Soap Fatty Acids (as	-~-			
above) 4-3	115			
Caustic Soda (37° Bé.) 6				
	lb.			
	lb.			
	10.			
No. 3				
Soft Soap Fatty Acids 12-15	lb.			
Hard Soap Fatty Acids 8-5	lb.			
Caustic Soda (37° Bé.) 12	lb.			
Water Glass (36–38° Bé.) 6	lb.			
Soda Ash 30	lb.			
	lb.			
No. 4				
Soft Soap Fatty Acids 18	lb.			
Hard Soap Fatty Acids (as	1.5.			
	lb.			
	lb.			
	lb.			
water diass (50-55 De.)	lb.			
Soda Ash 25	10.			

Soap Flakes

Water

27 lb.

To make high-grade soap flakes, a good quality charge consisting of 75% tallow and 25% coconut oil, with or without the addition of 2% or less of rosin, should be used. The mixture should be boiled and finished as for toilet soap, then chipped and dried. Care must be taken in drying in order to produce a uniform chip and avoid overdrying. The temperature of the soap chips should never fall below 30° C.; the temperature of the finished flakes should be between 40 and 45° C. The flakes should be milled twice to give transparency and polish. The most satisfactory shape to avoid breakage of very thin flakes is the square.

Soap for "Soap Noodles	,,	
	28	
Tallow or Hard Fat	4	g.
Caustic Soda (38° Bé.)	10	g.

Potassium Carbonate	
(30° Bé.)	10 g.
Water	10 g.
Salt Solution (24° Bé.)	10 g.
Sugar Solution (24° Bé.)	10 g.

Borax Soaps

Soap from Kettle	1000 lb.
Powdered Borax	130 lb.
Lye (40% Caustic Soda)	23 lb.
Perfume, etc.	sufficient

The soap is run into the crutcher, the borax, etc., added, and the whole crutched until the materials are thoroughly mixed. The physical condition of the soap is of less importance than when the soap has to cool in the frames and, therefore, the incorporation of larger quantities of

borax becomes feasible.

Various methods are available for the manufacture of soap powders, fillers being introduced before or after the soap is converted into powder. In the former case spoken of as the "continuous" process, the soda ash used takes up the excess water present with the soap, forming hydrated carbonate of soda and thus obviates the necessity of drying. A soap powder of this type suitable for laundry and general purposes can be obtained from the following formula:

Borax Soap Powder

Soap	42 lb.
Soda Ash	42 lb.
Powdered Borax	15 lb.
Salt	1 lb.

The soap is run hot from the kettle into the crutcher, and after thoroughly mixing with the soda ash and the borax, it is run over chilling rolls to chill the soap and crystallize the salts. The product is scraped off the rolls, the coarser particles being ground further. Alternatively, the mixture, after leaving the crutcher, is allowed to season for a few days, after which it is ready for powdering and packing.

Washing Powder

Fatty Acids	27.7-45.4 kg.
Sodium Perborate	4.8-13.5 kg.
Soda Ash	17.1-23.2 kg.
Water Glass	
(Dry Basis)	0.6-2.4 kg.

Abrasive Washing Powder

Soap		i, T	5 -10.2	kg.
Sodium	Carbonate		5.6-10	
Sand			73.7-81.5	

		Wa	ashing	Pow	$_{ m der}$
		\mathbf{F}	ormula	No.	1
Cut	into	small	pieces		

Hard	Soap	Waste	10	kg.
Dissolve Water			46	kg.
Add				

Water Glass	10	kg.
Sodium Carbonate, Calcined	39	kg.
Mix well to obtain homogene	ous	mass
No. 2		

Hard Soap Waste	20 kg.
Water	41 kg.
Water Glass	9 kg.
Sodium Carbonate, Calcined	35 kg.
No. 3	

Hard Soap Waste Water	40 kg. 38 kg.
Water Glass	4 kg.
Sodium Carbonate, Calcined	35 kg.

to get a 20% powder.

Note: the sodium carbonate is added only partially to the formulas 1, 2, 3, 3/3 is put on the bottom of the mixer before starting. Blow air into the warm mixture. Let cool for 24 hours.

Ammonia Washing Powder Hard Soap Powder (Alkaline) 1 lb. Ammonium Carbonate 1 lb.

Household Scourer

Colloidal Clay	1 lb.
Silica Floss	1 lb.
Alkaline Hard Soap Powder	4 lb.
Silicate or Carbonate of Soda	1 1h

Fermentative Washing Powder Sodium Carbonate 75 g. Bile, Precipitated on

Kieselguhr 25 g. 100 g. of this powder are applied to 50 kg. laundry batch.

Cold Processed Soap British Patent 403,500

A method for preparing "cold processed soap'' is to stir a mixture of 170 lb. of palm kernel oil with 9 gal. of 36° Bé. caustic soda solution. In a separate container, 6.5 gal. of a mixture containing equal parts of palm kernel oil and rosin is heated to 250° F., cooled to 110° F., and quickly added to the first mixture. After stirring for 10 seconds, the soap is run out through a valve in the bottom of the mixing pan, and sub-sequently treated in the usual manner.

Addition of rosin makes a more satisfactory and standard product than is usually obtained by cold process methods.

Cold-Process Carbolic Soap

For toilet purposes a cold or semiboiled soap is used, which retains the glycerin liberated from the fat. The following is a typical formula:

Formula No. 1

Coconut Oil	80 lb.
Tallow	40 lb.
Soda Lye (38° Bé.)	60 lb.
Phenol	3 lb.

The fat and lye are thoroughly stirred at 35° C. until combination occurs and the soap is streaky. The phenol (dissolved in a little water) is crutched well into the soap; perfuming is sometimes done with a little clove, lavender or rosemary oil. When cold the soap is cut into tablets and wrapped in air-tight package.

110. 2	
Bone Fat	150 lb.
Rosin	150 lb.
Carbolic Acid Solution	25 lb.
Caustic Soda Lye (37° Bé.)	150 lb.

The rosin and fat are melted together, and when the temperature is about 75° C. the carbolic acid is stirred in. The mixture is then added to the lye gradually, heating until the reaction is complete. The soap is framed and cooled and cut into bars of the usual size.

Cold Process Soap British Patent 432,227

Cold-process fat-resin soaps are made by treating fatty matter with just sufficient alkali for saponification, treating a mixture of rosin and fat or oil with alkali sufficient to saponify only the rosin, mixing the two products, and adding alkali to saponify the surplus fat. For example, 100 lb. of palm-kernel oil is stirred rapidly with 4.5 gal. of 36° Bé. caustic soda for 10–15 minutes, 4 gal. of a melt of rosin in an equal weight of palm-kernel oil is treated at 110–135° F. with 0.5 gal. of 36° Bé. caustic soda, the products are mixed, and immediately 1 gal. of 36° Bé. caustic soda is added, and the mixture stirred for a few seconds and run quickly into the frames, where it sets and saponification is completed.

Dry Cleaner's Soap British Patent 407,088

Soaps for use with dry-cleaning solvents, especially carbon tetrachloride or

trichloroethylene, consist of a fatty acid soap with a content of a polyglycol, with or without a chlorinated aliphatic hydrocarbon. An illustration is the following: 14.2 g. of sodium hydroxide is dissolved in 25 cc. of water and stirred into 100 g. of oleic acid and 100 cc. of trichloroethylene. Next 70 g. of triethylene glycol or 50 cc. of diethylene glycol is added. The product is dissolved in trichloroethylene.

Soaps Containing Pine Oil German Patent 616,029

Formula No. 1

a. Pine Oil Caustic Potash b. Coconut Oil Fatty	100 12.5	g. g.
b. Coconut Oil Fatty Acids	18-25	o.

Treat a at 80-100° C., neutralize the product with b.

No. 2

#10. #		
a. Pine Oil Caustic Soda (95%)	100	g.
a. Caustic Soda (95%)	4	ğ.
b. Fatty Acid	19	g.

As in No. 1. Solid, water-free soaps, high transparency.

Solid Pine Oil Soap U. S. Patent 2,007,974

Take one part water and two parts olive oil soap containing about 10% of water in the condition of flake or powder and when those are well blended stir in about one or two parts of pine oil. The vessel containing the mixture is placed in a kettle surrounded by glycerin and the temperature of the soap, water and oil is gradually raised to about 240° F. by heating the outer kettle. Steam is given off causing frothing of the soap with a great increase in volume of the mass. While some oils ordinarily begin to volatilize below this temperature, the soap raises the boiling point and permits them to be completely merged and held. When the heat, frothing and stirring have secured a uniform mixture, the mass is permitted to cool and solidify.

The solid soap lathers well, but slowly and yields at all dilutions a perfectly incorporated oil. It has the pleasant odor of pine oil but has the firm feel of anhydrous soap. The well fixed character of the oil is proved by the fact that the soap does not render white paper greasy after long contact with it.

Medicated Soaps

These types of soap can be made in two ways, either milled or by the cold process; as to their efficiency for the purpose for which they are intended, opinions differ, some claiming that they are of no value, others that certain complaints can only be cured by their use. Certainly much can be said for the latter statement, particularly when the complaint is in the nature of a skin disease such as eczema, and even without the addition of a specific body, toilet soaps which are superfatted with bodies such as lanolin or petroleum jelly naturally have a beneficial action on the skin.

No compound in skin soaps can compare with the well-known ichthyol variety. This compound can either be incorporated with flowers of sulphur and camphor or it may be used alone. Two mixings are given below containing these

bodies.

The first examples given are of the milled variety, which is certainly the best form of tablet both from appearance and as giving a perfect blend of the various bodies.

Ichthyol and Sulphur

,	-	
Soap Chips	28	lb.
Ichthyol	41/2	oz.
Vaseline	2	OZ.
Zinc Oxide	2	oz.
Flowers of Sulphur	2	oz.
Chlorophyll	$1\frac{1}{2}$	oz.
Medicated Perfume	4	oz.

Ichthyol

Soap Chips	28	lb.
Ichthyol	7	OZ.
Vaseline	2	0 Z .
Medicated Perfume	4	oz.
Zinc Oxide	2	0Z.
Chlorophyll	$1\frac{1}{2}$	oz.

The antiseptic value of the tablets is enhanced by the use of the medicated perfume, which gives the type of odor used in a well-known line on the market, having a ready sale as a medicated toilet soap.

Medicated Perfume

Eucalyptus Oil	18	cc.	
Terpineol	18	cc.	
French Lavender Spike Oil	18	cc.	
Red Thyme Oil	8	cc.	
Clove Oil	- 8	cc.	
Peru Balsam	6	cc.	
Camphor	3	o	

The soap and additions are milled in the ordinary way; it may be found necessary to mill more than the usual three times on account of the liquid nature of the additions. This may be obviated

somewhat by using the soap chips a little drier than the usual 76-77% fatty acids—say about 78-79%.

The chlorophyll used is the oil-soluble type, dissolved in a little medicinal paraffin, or if this is not available the perfume may be warmed slightly and used as medium.

All other kinds of medicated milled soaps can be made on the foregoing principle, leaving out the ichthyol, etc., and adding whatever is needed; the percentage used varies from 2½ to 5, the lower figure being more general.

The other variety is the well-known cold process soap, a very fine preparation for the feet. This soap, owing to the ease with which it is made, is one for the small manufacturer with his limited It contains permanganate of plant. potash, and the directions for its use are: Wash the feet and allow the lather to remain in contact with the skin a minute or so before rinsing. The instructions for its manufacture are as follows: Melt the tallow and coconut oil together, and at 120° F. pour in the caustic soda in a thin stream, stirring all the time; add the perfume and then the water, keeping the mass continuously on the move. When the soap is of the consistency of cream, which should be only about 3 to 4 minutes from the start. pour into a wooden frame and just crutch the permanganate solution here and there in the mass: do not thoroughly mix it in. The appearance obtained is similar to marble graining. After standing 45 hours, covered and free from draughts, the block of soap is ready for cutting, the size of tablets being usually

The mixing for the above soap is:

	Jour	χ,	•
Tallow	80		lb.
Coconut Oil	80		lb.
Caustic Soda, 66° Twaddell	80		lb.
Water	28		lb.
Perfume		1	lb.
Permanganate of Potash in			
1000 cc. Water	1/	4	lh.

Perfume

T CITUME		
Pine Oil	1	cc.
Cassia Oil	1/4	cc.
Lavender Spike Oil	1/2	cc.
Patchouli Oil	1/2	cc.
Ditolyl Methane	1/2	cc.

Another soap made as above, leaving out the permanganate and using in its place stavesacre seed oil with a different perfume, is also sold for the removal of head vermin in children, and may be included in the list of medicated soaps.

Perfume

Sassafras Oil	5 cc.
Geranium Oil	1 cc.
Sandalwood Oil W.I.	2 cc.
Terpineol	5 cc.

The active principles of the last-named soap are the stavesacre seed oil and the sassafras oil—a very effective combination. These few examples embrace the whole range of medicated soaps, the only alteration in other cases being the medicating substance, the percentage of which, as mentioned before, ranges between 2½ and 5.

Antiseptic Soaps

An odorless phenolated soap can be made by mixing in about 3% of a fatty acid phenol ester such as phenyl stear-These esters ate, palminate or oleate. are non-irritant to the skin and stable Iodine has been used in to alkalies. soaps. It does not have a very active antiseptic action when in the form of its compounds and is therefore employed as a solution in alcohol or in potassium iodide. Iodide is not stable however, as may be seen from the fact that soaps containing it change from brown to a light yellow in a short time. A better way of introducing iodine into soap is to add it in the form of a compound with an unsaturated acid such as oleic. A large number of so-called iodine soaps are made with potassium iodide and are quite stable, although they are not really iodine soaps.

Sulphur is a useful therapeutic for certain skin troubles. Its action is due to a mild antiseptic effect combined with reducing properties. Sublimed sulphur is generally used. The difficulty of getting sulphur into the water-soluble form may be overcome by using a combination of certain terpenes with alkaline sulphides and polysulphides. The solution of the clear brownish liquid in water gives a white emulsion with a slight alkaline reaction. It is non-irritant. A tar-sulphur soap is widely sold for the treatment of a variety of skin diseases. It is a brown soap prepared by dissolving 2 lb. of potassium sulphide in a small amount of water, and adding 20 lb. of yellow stock soap together with 4 lb. of birch tar oil. The mass is milled several times.

The manufacture of soap incorporating mercury or corrosive sublimate is not an easy matter. The mercury salt reacts rapidly with the soap to form complex insoluble compounds. An improved process for incorporating mercury makes the soap contain an excess of

free fatty acid, which prevents the chloride from reacting with the soap. In another process, the mercury salt is mixed with an alkaline casein solution, forming a mercury albuminate soluble in alkali.

Mercuric iodide is used in some soaps. It is best added by mixing 4 parts of mercuric iodide with 3 parts of potassium iodide and 2 parts of water, then incorporating the precipitated salt with the milled soap. The method of using nonionized complex mercury compounds is one that shows promise. These compounds give no black precipitate on addition of ammonium sulphide in the cold. Those which give no precipitate on prolonged standing are the best suited for the purpose.

Germicidal and Antisepti	e Soap
Coconut Oil Soap Base	50 g.
Cresol U.S.P. Mercuric Chloride	5 g.
1-2000 Solution)	45 g.
Iodine, Ichthyol, Campho	r Soaps
Formula No. 1	-
Soap Base	
Coconut Oil Ceylon	25 kg.
Caustic Soda (38° Bé.)	10 kg.
Caustic Potash (38° Bé.) Lanolin	2 kg. 1 kg.
Camphor	2 kg.
No. 2	8-
Iodine Soap	
Same, but add	
Potassium Iodide	1-1.5 kg.
in Water Hat	0 1
Water, Hot No. 3	2 kg.
Ichthyol Soap	
Same as No. 1, but add	
Ichthyol or	
Ammonium	
Ichthyolsulphate	1-1.5 kg.
Perfume	
Peruvian Balsam	120 g.
Lavender Oil	100 g.
Cassia Oil Benzoin, Tincture	100 g. 200 g.
Perfume only for No. 2 or	0
1 011 anie 0111 101 110. 2 01	710. 0.
Borie Acid Soap	
Doric Acid Soap	10

Sapamin-Phosphate (100%)

Triethanolamine Laurylsul-

Boric Acid

phonate

Distilled Water

Glycerin

10 oz.

5 oz.

5 oz. 20 oz.

60 oz.

Sand Soap	ence of rosin also assists. The propor-
Coconut or Palm Kernel Oil 20 kg.	tion of coconut oil is increased when the
Caustic Soda (38° Bé.) 11 kg.	soap is required to lather freely.
Pumice, Finely Powdered 10 kg.	
Solution of Benzoline,	Wool Scouring Bath
Tetralin 8 kg.	Olive Oil Soap 40 lb.
Turpentine Oil in Turkey	Ammonia 28% 20 lb.
Red Oil (1:1)	
Perfume 0.5 %	Transparent Glycerin Soaps
Mixture of	Formula No. 1
Lavender Spike Oil 5 cc.	
Rosemary Oil 4 cc.	a. Prepare a solution
Peppermint Oil 1 cc.	Caustic Soda (40° Bé.) 20 g.
Caraway Seed Oil 1 cc.	Alcohol (90–92%) 14 g.
Manufacture and Property and Commence and Co	Sugar 10 g. • Water 11 g.
Washing Tablets	Water 11 g. Glycerin 11 g.
Formula No. 1	Warm to 60-70° C.
Perborate of Soda 32 oz.	b. Add first melted
Granulated Borax 35 oz.	*
No. 2	Stearin, White 10 g.
Perborate of Soda 35 oz.	
Borax 17.5 oz.	Coconut Oil 18 g. Tallow, White 12 g.
No. 3	Castor Oil 4 g.
Perborate of Soda 27 oz.	
Borax 58 oz.	No. 2
No. 4	a. Caustic Soda (35° Bé.) 22 g.
Perborate of Soda 4 oz.	Alcohol 20 g.
Borax 12 oz.	Glycerin 20 g.
No. 5	Sugar 10 g.
Perborate of Soda 34 oz.	Water 10 g. Warm to 60-70° C.
Borax 18 oz.	
Soda Ash 22 oz.	b. Stearin 12 g.
In each of above formulas make up to	Coconut Oil 20 g. Castor Oil 5 g.
100 with soap. Crutch with soap; cut	•
into squares and dry.	No. 3
White the control of the state	English Transparent Soap
Wool Throwers Soap	a. Caustic Soda (38° Bé.) 50 g.
Olive Oil Foots 12 lb.	Alcohol (90-95%) 50 g.
Corn Oil 46 lb.	Sugar 17.5 g.
House Grease 20 lb.	Water, 60° C. 23 g.
Soda Lye, 36° Bé. 3 lb.	b. Pig Fat or Tallow 37.5 g.
Potassium Carbonate (Dry) 5¾ lb.	Rosin, Pale 12.5 g.
Potassium Hydrate (Solid) 23 lb.	Coconut Oil 50 g.
D	Filled (Chann) Transport Seems
Borax Laundry Soap	Filled (Cheap) Transparent Soaps
Finished Soap 1100 lb.	Formula No. 1 No. 2
Soda Ash 15 lb.	a. Caustic Soda
Solution of Carbonate of	(38° Bé.) 77 48 g.
Soda (30%) 25 lb.	Sugar 21 — g.
Solution of Metaborate of	Water 36 — cc.
Soda (s.g. 1-6) 25 lb. Silicate of Soda (40° Bé.) 85 lb.	Filling Solution * 90 50 cc. Alcohol 12 20 g.
Soap Stock 40 lb.	Alcohol 12 20 g. b. Coconut Oil 53.5 40 g.
	Pig Fat or Tallow 53.5 40 g.
The nature and proportions of the fats	Castor Oil 42 20 g.
and oils are important. In a general	* Filling Solution,
way the oils cottonseed, coconut, and palm-kernel, particularly the last two	Water, boiled 300 cc. 200
mentioned, take up and hold fillers better	Snger 51 g 70
than tallow and hardened oils. The pres-	Potassium Carbonate 52 g. 60 Salt 52 g. 40

Transparent Soap	
Hard Train Oil Fatty Acid	40 kg.
Soya Bean Oil Fatty Acid	60 kg.
Caustic Potash (50° Bé.)	42 kg.
Potassium Carbonate	13 kg.
Water	75 kg.

Filled Soap

$a. egin{cases} ext{Palm Kernel Oil} \ ext{Tallow} \ ext{Bone Fat} \end{cases}$	200 g.
a. { Tallow	100 g.
Bone Fat	100 g.
b. Water Glass	80 g.
$c. egin{cases} ext{Talc} \ ext{Water} \end{cases}$	60 g.
Water	60 cc.
d. Caustic Soda (25° Bé.)	370 cc.

Melt up a, keeping extra 20 of the palm kernel oil. Add b molten into kettle to d, and boil to right consistency. Add c as water-suspension. Now add salt water (23-24° Bé.) 8-10 cc., boil, test. If soap is too "sharp," add the remainder of the palm kernel oil until right. When tests show satisfactory results, boil 2 more hours and cool in covered kettle.

Soap Perfume

T. C. C.	
Cinnamic Alcohol	100 g.
Neroli	50 g.
Petitgrain (Grasse)	50 g.
Orangeflower Absolute	10 g.
Hydrarom Fleur d'Orange	5 g.
	15 %
Rose Otto (Bulgarian)	15 g.
Orris Concrete	5 g.
Costus (10%)	20 g.
Sandalwood, E.I.	80 g.
Bergamot	180 g.
Musk Ketone	40 g.
Musk Ambrette	20 g.
Coumarin	60 g.
Vetiverol	70 g.
Heliotropin	85 g.
Rhodinol, Pure	50 g.
Methylionone, Pure	60 g.
Benzoin Resinoid	60 g.
Phenylacetaldehyde (50%)	40 g.
2 (, , , ,	0

Automobile Tar Solvent

Naphtha	40	oz.
Ethylene Dichloride	90	oz.
Diglycol Laurate	5	oz.

Automobile Cleaner

Diglycol Laurate	10	fl.	oz.
Kerosene	2	pt.	
Naphtha	1	pt.	
Water	6	pt.	
Kieselguhr	1-2	ĺb.	

Bleaching Soda

a.	Water Glass, Commercial	
	(36–38° Bé.)	30 g.
b.	Water	25 g.
c.	Ammonium Carbonate	45 g.

Dilute a with b, warm up in a steamheated kettle with stirrer, add c and mix to homogeneous distribution. Pour hot on flat iron pans or on stone-floor, cool, turn with shovel, grind.

Stain Removing Powder U. S. Patent 2,022,262

For removal of iron stains from cotton and rayon textiles.

•	
Sodium Chlorite	1 oz.
Sodium Oxalate	1 oz.
Potassium Dihydrogen	
Phosphate	2 oz.

Dry Peroxide Bleaching Powder U. S. Patent 1,986,672

A bleaching powder comprises an apparently dry mixture obtainable by reacting a hydrogen peroxide solution with sodium bicarbonate and then adding anhydrous sodium carbonate all in the proportions of substantially 10 parts of 30 volume per cent of hydrogen peroxide, 6 parts of sodium bicarbonate and 135 parts of anhydrous sodium carbonate.

Bleaching and Washing Powder French Patent 783,871

Formula No. 1

10 kg.

49 kg.

Sodium Perborate

Soap

Sodium Pyrophosphate Soda Ash Magnesium Silicate	14 kg. 8 kg. 1 kg.
No. 2	
Sodium Perborate	15 kg.
Sodium Hexametaphosphate	10 kg.
Soda Ash	9 kg.
Magnesium Silicate	1 kg.

Stone, Brick and Masonry Cleaner U. S. Patent 1,990,383

Forty gallons of soap-bark extract formed from 9.5 lb. of soap-tree bark by steeping in water are mixed with rosin oil 1.25, raw linseed oil 1.25, an aqueous gum tragacanth solution (containing 1.25 oz. of the gum), (114 to 22%) hydrochloric acid 10 gal.

Brick and Masonry Cleaner

Use a saturated water solution of ammonium bifluoride.

Drain Cleaner

Caustic Soda, Powdered	15	oz.
Chalk, Powdered	25	oz.
Caustic Potash, Powdered	60	oz.
Keep dry and pack in air-t	ight	tins

Washing Compounds for Use in Canning

The greatest surface is cleaned by a solution of a mixture of sodium hydroxide 2.8, soap 0.2, water glass 14.1 and sodium hypochlorite 4.8 (chlorine 2.3%) but this has some corrosive action.

Cleanser for House Façades

Trisodium Phosphate	75 g.
Sodium Metaphosphate	20 g.
Turkey Red Öil	3 g.
Sodium Hydroxide	2 g.
Water to desired	concentration

Floor Bleaches

Oxalic acid has long been used to bleach or whiten discolored wood in its natural finish, especially floors. After applying this chemical, however, the wood is left so white that the spot usually must be stained lightly to restore it to the shade of the surrounding wood. Sodium perborate, which is sold in drug stores for use as a mouth rinse and a tooth powder, is a far milder bleaching agent. Although one may have to rub the moistened powder on the discolora-tion a longer time than if an oxalic acid solution were used, the after effects are not so conspicuous. It is also particularly effective when mixed with equal parts of sodium metasilicate.

Cleanser for "Parquet" Floor

	ntv

Caustic Soda (128-130°)	6.64 kg.	
Water	26.36 kg.	
Red Oil (Oleic Acid)	45.45 kg.	

A 44.

and a			
Alcohol,	Denatured	45.4	l.

The whole poured into

Tric	nioroetny	lene			900	kg.
The	product	gives	a	stable	em	ulsio

with water.

Cleansing Preparation for Galoshes

	Carnauba Beeswax	Wax,	Fat	Gray	1	kg.
u.	Beeswax				0.5	kg.

1	Olive Oil Soap Borax Capillary Syrup		kg.
. 1	Borax	0.5	kg.
b. {	Capillary Syrup	0.3	kg.
	Water	25	l.

Melt up a, dissolve b by short boiling, add b to a and stir until cooled, then add

Thinner (as above)	12	1.
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Cleanser for Dishes, Glasses, etc.

Formula No. 1

Trisodium Phosphate	45 g.
Sodium Metaphosphate	53 g.
Caustic Soda	2 g.
No. 2	7
Trisodium Phosphate	55 g.
Sodium Metaphosphate	43 g.
Caustic Soda	2 g.
No. 3	
Trisodium Phosphate	75 g.
Sodium Metaphosphate	23 g.
Caustic Soda	2 g.
No. 4	

Trisodium Phosphate

(Monohydrate)	15 g.
Sodium Metasilicate	40
(Pentahydrate)	40 g.
Sodium Metaphosphate Caustic Soda	40 g. 5 g.

Mechanical Dishwashing Preparation Sodium Metaphosphate 40 oz. Trisodium Phosphate 15 oz. Sodium Silicate 40 oz. Sodium Hydroxide 5 oz.

Glass Cleaners

Glass Cleaner in Cake Form

Infusorial Earth, Finest		
Powder	4	OZ.
Precipitated Chalk	2	oz.
White Soap	- 2	oz.

Boiling Water Reduce the soap to fine shavings and dissolve in the boiling water. Then add powders which have been previously mixed and put through a fine sieve. Press into molds the size of the cake required and allow to dry.

L			
	White Soap	750	g.
	Sodium Carbonate	20	OZ.
	Hot Water	120	cc.
	Infusorial Earth	250	o.

Dissolve the soap (in fine shavings) in the hot water in which the sodium salt has been dissolved. Then add the infusorial earth in very fine powder. These soaps may be perfumed slightly by the addition of equal parts of oil of sassa-

fras	and	cedar	oil	to	sui	t.	Th	ese
		very 1						
		g to i						
the p	ropei	ty of	abso	rbin	g co	nsi	dera	ble
water								

The following formula is another

example:

Powdered Pumice Stone		2	oz.
Ammonium Oleate		-	oz.
Ammonia (28%) to	make	16	oz.
Shake before using.			

Cleaning Mixture for Beer Glasses Use 1-3 g. per l. water of one of the mixtures (finely ground):

Formula No. 1

200	
350	g.
50	
700	g.
300	g.
800	g.
200	g.
150	g.
850	g.
	700 800 800 200

Window Glass Cleaner

	2 **
a.	MIx

ъ.

Neuburger Chalk, Ppt.,	
Finest	40
Viennese Lime	20
Calcium Carbonate, Ppt.,	
Heavy	25
Bolus, White	15
And grind with a mixture of	
Water	90%
Alcohol, Denatured	5%
Ammonia (sp. g. 0.91)	5%

Gun Cleaner and Solvent

Turpentine	2	fl.	OZ.
Methyl Acetone	1	fl.	oz.
Sperm Oil	2	fl.	oz.
Butyl "Cellosolve"			oz.
Kerosene	4	fl.	oz.
Lanolin	1	oz.	

Special Cleanser for Very Dirty Hands Coconut or Palm Kernel Oil

Coconic of Faim Refuel On	
Fatty Acids	6 g.
Soya Bean, Linseed, Peanut	
Oil Fatty Acids	6 g.
Castor Oil Fatty Acid	3 g.
Pine Oil	6 g.
Alcohol	6 g.
Lanolin	1 g.

Caustic Potash (50° Bé.) 6 cc.
Water 6 cc.
Pumice, Fine Powder until pasty
Citronella, ''Spike'' Oil,
Terpineol as Perfume to suit

Antiseptic Cleaner for Ice Cream Freezers

At the conclusion of the freezing operation drain the ice cream from the freezer. Rinse the strainer, hopper, and outside of the freezer, particularly at the head, with cold water. Fill the freezer two-thirds full of cold water, run one-half minute, and drain.

Fill the hopper full of water at 140° to 145° F. and add a half pound (1 cup full) of cleansing powder. Wash the strainer, hopper, and outside of the freezer with a brush. Drain the solution into the freezer (the freezer should be at least two-thirds full), run one-half minute, and drain the freezer.

Remove the head, scrub with a brush, being certain to clean out the front bearing. Wash the bearing end of the dasher with a brush, remove from freezer and wash. Place dasher and head in sanitary

place until used.

Before using the freezer, fill the hopper with water at 100° to 110° F., making certain that the screen is covered. Add sufficient chlorine to give 100 p.p.m. and stir well. If desired, the chlorine solution can be pumped into the hopper from a special tank. Pour some of the chlorine solution into the front bearing. Place dasher in freezer and fasten the head in place. Drain the chlorine solution into the freezer, operate the freezer one-half minute, and drain. The freezer is then in excellent sanitary condition, except that the rear bearing may be contaminated, and is ready for use.

Lavatory Cleaner

One method is to add niter cake (acid sodium sulphate) to the water in the bowl. Another consists of a mixture of sodium carbonate (16 parts) and caustic soda (3 parts), and there are others depending on the liberation of chlorine.

A cleaner can be made up of sodium sulphate (88 parts), sulphuric acid (9 parts), and diatomaceous earth or some other fine abrasive material (3 parts).

Another suggestion is to mix powdered soap with four times its weight of powdered potassium carbonate.

Coconut Oil	10	lb.
Potassium Hydroxide	1	lb.
Sodium Hydroxide	1	lb.
Water	10	lb.

Dissolve the potassium hydroxide and sodium hydroxide in the water and mix with the coconut oil. Set aside in a warm place for a few hours to saponify. Test for neutrality and dissolve the product in 6 oz. of water. The resulting liquid soap does not cake and lathers freely when used in small quantities.

Laundry Bleach

Chlorinated Washing Soo Water		11/2	lb. lb. gal.	
		-		

Allow to stand for a few days and filter.

Laundry Blue Good Quality

Formula No. 1

Ultramarine	60 lb.
Bicarbonate of Soda	40 lb.
Glucose	12 lb.
No 9	

Cheen Quelity

Oneap withing	
Ultramarine	18 lb.
Kiln-Dried Blue Earth	20 lb.
Terra Alba	15 lb.
Bicarbonate of Soda	45 lb.
Glucose	10 lb.
No. 3	
Lime	5 oz.

Water 10 oz.

Stir until smooth and mix with a hot solution of

OLLIOTOTE OF			
Dextrin, Yellow	5	oz.	
Water	3	oz.	
Glycerin	5	oz.	
Phenol	0.2	OZ.	
Ultramarine Blue Powder	75	oz.	

Ultramarine Blue Paste, Laundry

- 1 0140	·	UZA.
a. Water	10	oz.
Soak cold, then warm to	dis	solve.
Yellow Dextrin	5	oz.
b. Water Glycerin (sp. g. 1.23)	3	OZ.
Glycerin (sp. g. 1.23)	5	oz.

Mix both parts warm, conserve with 0.2% nipagin, moldex or phenol, etc., and grind now with

Ultramarine Blue or

Imitation of Ultramarine 75 oz. formed by precipitating anilinlakes (dyestuff) on insoluble inorganic bodies on china clay or white bolus.

Laundry Sour U. S. Patent 1,998,819

A souring composition is formed of sodium fluosilicate 84, sodium acid fluoride 15 and gelatin 1, all parts by weight, or the like.

Cleanser	for	Hunting	Calf	Leat	her
Trioxyme	ethyl	ene		70	g.
Cleaning	Ber	zoline		30	cc.
Oxalic A				5	g.
Liquid S	oap				cc.
Mix thor	ough	lv.			

Cleanser for Sporting Leathers

1 0	
Water	75 cc.
Acetic Acid (80%)	5 cc.
Alcohol (95%), Denatured	. 30 cc.

Cleaner and Disinfectant for Metal Articles

U. S. Patent 1,937,229

Sodium Silicate (D. 1.38) plus 500 g. of following	300 g.
Sodium Hypochlorite	
(D. 1.125)	562 g.
Caustic Sodá (D. 1.383)	250 g.

Bleach-Bath for Used Oil Corks (e.g. of Olive Oil Bottles)

A ¼ to 2% solution of above is used.

a. Remove fats with hot alkaline solutions, as soap, soda, trisodium phosphate; wash thoroughly with hot water.

b. Hydrogen Peroxide (1.5-1.6%) 10 l. Ammonia (25%) 200 g.

Treat corks cold (18-20° C.) for about five days, adding every 8 hours new
Ammonia (25%) 40-50 g.

Oven Cleanser Formula No. 1

Olein, Distilled	40	oz.
Stearin	10	oz.
Mix warm.		
Spindle Oil	40	oz.
Tetralin	9	oz.
Ammonia (sp. g. 0.91)	1	oz.
Emery or Pumice or Tripoli		
sufficient to make	pa	sty

No. 2

Ceresin	(56-58°	C.)	7	g.
Olein			17	g.
Mineral	Oil		6	g.

Slate Powder	about	10	g.
Chromium Oxide		15	
Carborundum or Emery	about	45	g.

Printing Form and Cylinder Cleaner

Test Benzoline	00
(B. P. 130–150° C.)	80 cc.
Xylol	15 cc.
Petroleum Oil	5 cc.
Ignition point should be	e over 21° C.

Rug Cleaner

Coconut Oil Soap Ammonia (28%) Glycerin Water	12 2.8 7.9 77.3	oz.
water	11.5	oz.

Radiator Cleaner

Compound for use in hot force pump automobile radiator flushing tanks.

76% Flake Caustic Soda	60 lb.
Sal Soda	30 lb.
Rosin	10 lb.
Use about 40 lb. to 75 gal	. water.

Dry Cleanser for Wallpapers

Wheat Starch 35 oz. Sodium Chloride, Saturated Solution 65 oz.

Warm upon water-bath and stir until sufficiently plastic. Shortly before the end of this treatment, when cooled, add a little naphtha. Apply like a sponge eraser. Pack in air-tight tins.

Wall Cleaner

Corn Flour	90 lb.
Copper Sulphate	9 lb.
Alum	1 lb.
Mix and dissolve in boiling	water.

Scouring Soaps

The following is a soap-sand cleaning preparation that has a wide sale for household and general purposes. It takes the form of a palm oil and coconut oil soap, which is then liquored down in the same pan with carbonate of potash, carbonate of soda crystals, silicate of soda 100° Twaddell, and water.

Melt the two oils, pass in steam, and then pour in caustic soda gently, adding a little water from time to time to keep the soap smooth. Saponification will proceed fairly easily, as the palm oil soon takes up. When all the caustic soda has been added, pour in the remainder of the water in such a way that the mass never ceases to simmer; the operation should

take about 4 hours. Towards the end add the other ingredients, which will dissolve easily, as the finished product is very similar to a liquid soap.

Let the soap liquid cool to about 90° F., and to 10 lb. of dried common sand add the same amount of the above soap. All the time the soap is being added, the mass must be stirred rapidly, and when it resembles a thick sludge it will be ready to pour into tins. The only precaution to take is that the mass must not be poured in too warm, as naturally the sand would precipitate in the tins; this part of the operation can only be perfected by actual experience and must always be done very carefully, but no difficulty should present itself if all directions are carried out as given.

Mixing

Coconut Oil	4	lb.	
Red Palm Oil	69	lb.	
Caustic Soda, 60° Twaddle	37	lb.	
Additions			
Carbonate of Potash	5	lb.	
Soda, Sal	15	lb.	
Silicate of Soda,			
100° Twaddle	21	lb.	
Cresylic Acid	3	lb.	
Pine Oil	11/2	lb.	
Orange II. (Color)	1/4	oz.	

The whole mass of soap and additions should total up to 784 lb., with the addition of water.

A hand-cleansing soft soap can be obtained by the use of a carbolic soft soap, preferably one made from vegetable and not fish oils, using the same proportions of soap and sand as in the previous example, but it would be better in this case to use, in place of the sand, pumice powder of 120 mesh. Sand is, naturally, coarse and cheap; better scouring agents might be used, such as silver sand, or pumice powder of 60, 90, or 120 mesh, according to the nature of the finished article desired.

Scouring Powder

Silica 100–125 mesh	75 oz.
Soda Ash	13 oz.
Trisodium Phosphate	8 oz.
Soan Powder	4 oz.

These materials in powdered form are thoroughly mixed together and are ready for use as such.

Stain Emulsifier

Liquid Soap (15%)	40 cc.
Liquid Soap (15%) Turkey Red Oil (100%)	25 cc.
Decalin	4 cc.

Turpentine	4 cc.
Ethylene Glycol	10 ec.
Methylene Glycol	10 ec.
Methanol	5 cc.
Terpineol	2 cc.
-	

Removing Glue Stains from Wood

Casein and vegetable glue stains can be almost entirely removed by sponging the stained surface with an oxalic acid solution prepared by dissolving 1 oz. of oxalic acid crystals in about 12 oz. of water. Still better results may be obtained by moistening the wood first with a sodium sulphate solution made up in the same concentration as the oxalic acid. In this way stains have been almost eliminated.

Remover	for	Tobacco	Stains	on F	'ingers
Hard S	Soap	Powder		40	oz.
Marble				20	oz.
Alcohol	l, De	enatured		40	oz.

Soap hands with this mixture, rub at the same time with finest pumice powder.

Removing Pitch or Varnish from Hands or Glass

Household Scouring Powder Dutch Cleanser type Acetone

ne

sufficient to make a thin paste

Rub the hands or article to be cleansed with this paste. The viscous impurity is at once dissolved in the acetone, and is absorbed into the powder mass. Within a minute or two the acetone evaporates, leaving a mealy or dry powder which can be dusted off, or in suitable cases as with the hands, washed off. Do not use on a painted, varnished or lacquered surface, which would be injured by the acetone. This is a very economical material for the purpose.

Soot Destroyer

Salt	•	85	oz.
Copper Sulphate		8	oz.
Zinc Dust		7	oz.

Steamship Chimney Soap

Soft Soap, Brow	vn 20	g.
Water	12-15	cc.
Potassium Carbo		g.
Hexahydro-cresol		cc.
Decahydro-napht		cc.
Sodium-Di-Isobu	tyl-naphtha-	
lene Sulphonat	te 1.5–2	g.

Cleanser for Lampblack-Dirtied Surfaces

a. Olein or Oil Fatty Acid 45.45 kg.
 Caustic Soda

b. { (128-130°) 6.64 kg. (128-130°) 6.64 kg. c. Alcohol 45.4 l.

Saponify a with b on water bath, dissolve, then warm (below 70° C.) in c. Add stirring

d. Tripoli 900 kg. and thin 10 times with water.

Floor Sweeping Compound Formula No. 1

Sawdust, Dyed Green with Aniline Dye, e.g., Brilliant

Green 35 kg.
Rock Salt 35–40 kg.
Mineral Oil, Deodorized
(2–3° E. at 50° C.) 25 kg.

No. 2

The following is a representative formula for floor sweeping compounds.

, coping	compour	us.
	10	lb.
	32	oz.
	2	oz.
	8	oz.
	4	lb.
	recping	32 2 8

Tinned Ware Cleaner

Sodium carbonate alone is not a satisfactory cleanser for milk containers of tinned copper, since it slowly removes tin as stannite owing to the presence of dissolved oxygen. The exposed copper produces an ''off flavor'' in the milk. The addition of sodium sulphite reduces the rate of attack to nearly 0.1. It is much more effective than a number of other reducing agents tried because it is more active in reducing the amount of dissolved oxygen. Suitable proportions are 1 lb. sodium sulphite and 10 lb. washing soda, 1 lb. sodium sulphite and 4 lb. sodium hydroxide (or sodium carbonate).

Type Cleaner

J.			
Butyl "Cellosolve"	. 1	pt	•
Diglycol Laurate		fl.	

Cleanser for Velvet Shoes

Water	100	cc.
Potassium Alum	1	g.
Alcohol	20	čc.
Turkey Red Oil	5	cc.

Composition for Cleaning Walls, Paint, etc.

French Patent 774,876

The composition contains corn flour 455, copper sulphate 40, alum 5 parts and is mixed with boiling water for use.

Painted Woodwork Cleaner

This specialty product quickly removes dirt from paint and leaves the painted surface with a bright, clean, lustrous finish. The diglycol stearate serves the combined purpose of emulsifying the dirt as fast as it is dissolved and of imparting a lasting natural luster to the cleaned surface. The product, therefore, may truly be said to both clean and shine in one operation. This new type of cleaner is made to the following formula.

Diglycol Stearate		lb.
Kerosene		gal.
Trisodium Phosphate	$4\frac{1}{2}$	OZ.
Water	12	pt.

Method of manufacture: The diglycol stearate and kerosene are heated together in a double boiler until the wax is thoroughly dissolved. Kerosene is inflammable, therefore care should be taken to prevent it from catching on fire. The trisodium phosphate is dissolved in the water and heated in another container to a temperature of about 150° F. The hot water solution is then added to the hot kerosene solution while stirring at high speed. Stirring should be continued at a good rate until the mixture is of even milky consistency. Mixing may then be continued at a slow rate until the batch has cooled to around 85° F.

This product is applied in the usual manner by rubbing with a rag or cloth. The same product may also be used for cleaning automobiles before waxing. However, for this service 12 oz. of fuller's earth should be thoroughly worked into the above batch after it has cooled over night. The fuller's earth should not be added until cooling is complete. With this addition a product is produced which cleans rapidly and with-

out scratching the finish.

"Soluble" Pine Oil Fluid

A satisfactory clear, pale straw pine concentrate, which is perfectly stable and gives a dense milky emulsion when added to water can be made from the following formula:

Heavy White I	Pine Oil	70	cc.
Oleic Acid		12	cc.
Water		18	cc.

The procedure is very simple—dissolve the oleic acid in the pine oil in the cold, and neutralize carefully with a 28% solution of caustic potash or soda. Caustic potash gives a slightly better color than caustic soda. By this method no heat whatever is required.

Soap Towel U. S. Patent 1,969,900

A towel for cleaning surfaces consists of a paper towel carrying a detergent composition including pine oil about 3-10 parts, a soap about 0.3-0.6 parts and water about 85-95 parts.

Sodium Metasilicate Solutions

Solutions containing 20 g. per l. of a commercial detergent preparation (sodium silicate 40, baking soda 30, soap powder 20, sodium perborate 10) show turbidity a few hours after preparation followed by precipitation; this renders it useless. Solutions of 5–10 g. per l. of sodium silicate begin to precipitate in presence of 35–40 g. baking soda per l. and precipitation is instantaneous with more than 40 g.; a solution of 15–30 g. per l. of sodium silicate begins precipitating in presence of 20–25 g. baking soda. Substitution of trisodium phosphate for baking soda immediately corrects the trouble.

Movie Film Cleaner

Carbon Tetrachloride 65 oz. Ethylene Dichloride 10 oz. Petroleum Ether 25 oz.

This composition is used to clean dirt, greasy spots and all foreign matter off of both faces of a movie film without affecting or having any solvent action on the film or gelatin coating itself.

The petroleum ether is a light fraction distillate with an end point under 100° C. These solvents are mixed together and

are then ready for use.

Benzine Soap

Dissolve 10 lb. of curd soap in boiling water, add a strong solution of magnesium sulphate slowly with stirring until it is all transformed into an insoluble mass, skim off the magnesium soap thus formed and purify by boiling it with fresh water. Remove the excess of moisture by squeezing through a cloth and pressing. Place the soap in a jacketed copper kettle and heat slowly to 266° F, turn off the heat and add 7 lb. of deodorized petroleum distillate. Dissolve

the product in 22 gal. benzine. If the solution is not clear the water has not been completely removed. For garment cleaning use 1 qt. of this solution for 25 gal. of benzine.

Dry Cleaning Solvents for "Celanese"

The following chemicals are safe for cellulose acetate fabrics: gasoline, Stoddard's solvent, cleaner's naphtha, kerosene, dilute alkalies (such as soap and water, soda, ammonia, sodium hypochlorites, Javelle water and washing sodas), glycerin, carbon disulphide, turpentine, all the hydrosulphite solutions (such as decolorite, blanket, sulphogen, burmol, paragene and lykopon), petroleum ether, vaseline, toluol, xylol, good grades of wood or denatured alcohol used cold and washed thoroughly, sulphuric ether, trichloroethylene, benzol, which is one of the best all around spotting chemicals, and unadulterated carbon tetrachloride, which is rapidly taking the place of chloroform. It is a known fact that carbon tetrachloride will absorb a small amount of moisture from the air if the container is left open. If moisture is present this powerful solvent is crippled and will not be as effective as when dry. To test carbon tetrachloride for purity, take two parts mineral oil, such as Nujol, and one part carbon tetrachloride. Mix. If this mixture becomes milky it denotes the presence of water in the carbon tetrachloride and in this condition should not be used for spotting purposes.

Dry Cleaning Soap

Curd Soap	3	0 oz.
Water		0 oz.
Ox Gall (Dried)	1	0 oz.
Soda Ash		5 oz.

Shred the soap and dissolve in hot water, adding the ox gall and soda. Evaporate the solution until on cooling, a sample on a slab sets solid. Pour the mixture into trays or molds. The disadvantage of such a preparation is its rather unpleasant smell.

Dry Cleaning Soap British Patent 407,088

Fourteen and two-tenths grams of sodium hydroxide is dissolved in 25 cc. water and stirred into 100 g. oleic acid and 100 cc. trichloroethylene; 70 g. triethylene glycol or 50 cc. diethylene glycol is added and the product is dissolved in trichloroethylene for dry cleaning.

Textile Soap French Patent 658,412

A		•		
Castile Soap			200	lb.
Tallow Soap,	Powdere	$^{\mathrm{d}}$	95	lb.
Soda Ash			20	lb.
Borax			10	lb.
Turpentine			25	lb.
Caustic Alkali			20	lb.
	lissolved	in 30	of wa	ter

Kier Soan

zzrez word		
Red Oil	2050	lb.
Rosin	1050	
Soda Ash	290	lb.
Caustic Soda (50° Bé.)	746	lb.
Water to make	11000	lb.

Ox Gall Soap

Since ox gall derived from bile has an unpleasant smell, an improved method is to add to soap solution about 14% of sodium cholate, the sodium salt of cholic acid which is a purified decomposition product of bile. It is claimed thus that the advantages of the detergent power of ox gall are obtained without the accompanying odor.

Rose Soap

a. White Tallow Soap	10000 kg.
b. Moistened Cinnabar	60-80 kg.
Rose Essence Clove Essence	40 kg.
Clove Essence	15 kg.
c. Cinnamon Essence	10 kg.
c. Cinnamon Essence Neroli Essence Bergamot Essence	10 kg.
Bergamot Essence	30 kg.
Perfume	_

Windsor Soap

10000 kg.

60 kg.

15 kg.

a. White Tallow Soap

r Bergamot Essence

Caraway Essence	25 kg.
Clove Essence	16 kg.
$b. \begin{cases} \text{Caraway Essence} \\ \text{Clove Essence} \\ \text{Thyme Essence} \end{cases}$	25 kg.
Perfume	
or	
Bergamot Essence	25 kg.
Caraway Essence	60 kg.
b. Caraway Essence Rosemary Essence	15 kg.

Witch Hazel Soap

Fine Lavender Essence

Witch Hazel Extract U.S.P.	10	oz.
Distilled Water	10	oz.
Triethanolaminelauryl-		
sulphonate	80	oz.

Perspiration Odor Destroying Soap Aluminum Chloride Crystals 3 oz. Hydrochloric Acid $\frac{1}{10}$

Normal Triethanolaminelaurylsulphonate 1-2 oz.

96-95 oz.

Soft Soap Manufacture

Soft soap contains normally 40 to 44% of fatty acids. The best method of saponification is to take the calculated quantities of alkali sufficient to effect complete saponification, with an excess of 1 to 1.5% alkali. The caustic solution, preferably of a density of 30° Tw. (about 19° Bé.), is brought to a boil and the melted charge added as quickly as possible without the contents frothing over. Emulsification follows with rapid The process is usually saponification. complete in a few hours' time, water being added when necessary. If rosin is to be incorporated, it is best added after the other stocks have been saponified.

Unless castor oil is present a soft soap charge cannot be worked with caustic soda alone. With caustic soda, castor oil will form a soft soap. Soft soaps can be made with castor oil in which varying proportions of other stocks have been introduced with the substitution of varying proportions of caustic potash for caustic soda. Saturated fatty acids tend to give stringy soap even with potash. The higher these are in the homologous series, the more pronounced is the stringiness.

The percentage of caustic soda which can be substituted for caustic potash will depend on the percentage of castor oil introduced into the blend. Practical experiments indicate that about 2% of caustic soda can be substituted for caustic potash for every 1% of castor oil introduced into the blend. Use of caustic soda in this way does not affect translucency and gloss.

Linseed-soda soap is stringy, but the corresponding potash soap is non-stringy with a desirable body. Peanut oil-soda soap is stringy but the potash soap is not. The following blends suggest the possibilities for soft soap manufacture:

The charges given below produce a stringy soap with 80% caustic potash and 20% equivalent caustic soda, but have the right non-stringy body with caustic potash only:

Formula No. 1

Peanut oil 20 parts, linseed oil 50, cottonseed oil 20, tallow 5 and rosin 5.

No. 2

Peanut oil 20 parts, linseed oil 50, cottonseed oil 20 and tallow 10.

No. 3

Linseed oil 60 parts, cottonseed oil 30 and rosin 10.

No. 4

Linseed oil 65 parts, cottonseed oil 25, rosin 10.

No. 5

Linseed oil 67 parts, peanut oil 13, cottonseed oil 10, tallow 5 and rosin 5 can be used with 80% of caustic potash and 20% equivalent caustic soda to give an almost non-stringy soap with only a slight thready tendency.

No. 6

Linseed oil 73 parts, cottonseed oil 15, rosin 10 and coconut oil 2 can be used with 70% of caustic potash and 30% equivalent caustic soda to give a nonstringy soap. In general it is preferable to use more potash. This represents the lower limit of potash with this type of blend.

No. 7

Linseed oil 73, castor oil 20, rosin 5 and coconut oil 2 gives a correct nonstringy soap with 60% caustic potash and 40% equivalent caustic soda, due to the introduction of eastor oil.

The following blends with higher percentages of castor oil give non-stringy soap with caustic soda alone:

No. 8

Linseed oil 38, castor oil 50, coconut oil 2 and rosin 10 parts.

No. 9

Linseed oil 32, castor oil 45, coconut oil 3 and rosin 20 parts.

No. 10

Linseed oil 50, castor oil 35, coconut oil 3 and rosin 12 parts.

Soap Rancidity, Preventing

This is best done by kneading into the dry soap, before milling, .7% of the following mixture:

Beeswax 300, anhydrous lanolin 400, liquid paraffin 390, water 300, borax 17, sodium thiosulphate 690, water 200. Melt together the wax, lanolin and paraffin oil; then dissolve the borax in 300 parts of water and pour this solution in a thin jet into the hot mass of molten fats at a temperature of about 95° C. Boil for a few minutes longer, then set aside and let cool to 50°, stirring frequently. Pour the hot solution of sodium thiosulphate in 200 g. of water into the fat-borax emulsion in a thin jet and stir until smooth. In some cases, for example

when using an unusually large quantity of perfume, it is advisable to add 1% of the following:

Beeswax 200, anhydrous lanolin 600, liquid paraffin 390, water 200, borax 17, sodium thiosulphate 690, water 200, sodium silicate 450, granulated sugar 253.

Superfatting Soap

Use of a superfatting agent undoubtedly improves the texture of soap, making it more plastic and easily worked. It also tends to neutralize any alkali which might be present, and thus remove harshness which might irritate sensitive skins. A good mixture for this purpose consists of equal parts of stearin and white petroleum jelly, or 2 parts stearin, 1 part lanolin, and 1 part white petro-

leum jelly. These are inelted, mixed, allowed to cool, and 1 to 1½ lb. added per 100 lb. of chips added with the other ingredients at the mixing stage.

Soap Spirit

Olive Oil		1000	ec.
Caustic Potash	(50%)	about 396	ec.
Distilled Water		2600	cc.
Alcohol (90%)		6000	ec.

Softener for Hard Water

Soldend for fitting water
Water Glass (36–38° Bé.) 25 oz.
Water 25 oz.
Ammonium Carbonate about 50 oz.
Mix well (warming), pour off to
solidify the paste. When cool, grind and
add to 95 oz. of the material.

50 oz.

Trisodium Phosphate

TEXTILES, FIBERS

Starches and Siz	zes for mula N		Shee	ting
Cornstarch Castor Oil Color Water Boil together	until si	220–2	00 lb. ½ pt to su 40 ga	it
1,011 108001101	No. 2			
Cornstarch Gypsum Castor Oil Color Water	No. 3		00 lb. 80 lb. 1 pt. to su 40 ga	it
Cornstarch	110. 5		60 lb.	
Lard			5 lb.	
Blue Dye		2	-4 oz.	
Water		1	20 ga	1.
	No. 4			
Cornstarch China Clay Lard Color Water	No. 5	1	65 11 10 11 5 11 to sui 20 ga	o. o. it
D 1 1 01 1	140. 9		00 11	
Potato Starch Steeped Flour Slaked Lime China Clay Elaine (Red) (Blue Color Water to make Boil for 1–2 m	Oil e	Tw.)	00 lb. 10 ga 15 ga 15 ga 3 pt. 12 oz. 20 ga	l. l. l.

	Cream Sizing	
	Tallow	36 lb.
	Calcium Chloride	6 lb.
	Starch	7 lb.
	Gum Arabic	6 lb.
	Water	45 lb.
	Cook and stir at 220-230°	F. for 1-
1	nours.	

Sizing Rayon and Silk French Patent 779,584

Rayon and silk are sized to give firmness, elasticity and suppleness by a solution in water of

	Formula No.	1
Stearic	Acid	15 g.
Glue		35 g.

Gum Arabic Soap Glycol Stearate Borax Pepsin No. 2	8 g. 32 g. 18 g. 2 g. 0.15 g.
	0.0
Lauric Acid	20 g.
Gelatin	34 g.
Soap	30 g.
Gum Arabic	
Ethylene Glycol	
	6 g.
Borax	2 g.
$\mathbf{Trypsin}$.05 g.
No. 3	
Glue	20 g.
Gum Arabic	0
Glycol Stearate	
Soap	18 g.
Glycerin	8 g.
Borax	1.2 g.
Pepsin	.01 g.
	8.
Rayon Size	
Calcium Resinate	20 lb.
No. 1 Lard Oil	
	10 lb.
Xylol	35 lb.
Damar Gum	10 lb.

Manipulation: Dissolve the damar gum in the xylol and add the other ingredients at 50° C. Then cool slowly with agitation.

Light Goods Sizing Formula No. 1

Soluble Potato Starch $1\frac{1}{2}-2\frac{1}{2}$ lb. Glucose 3-5 pt. Water 5 gal.

The starch and glucose are entered into the water and the whole brought to a boil and continued at that temperature until the starch particles are entirely cooked, which will depend upon the particular type of starch used. Before using, the mixture should be allowed to cool to a temperature of about 180° F. The purpose of the glucose is to impart a soft feel to the material and may be omitted.

No. 2

Another mixture that is suitable for setting goods other than those constructed of rayon, is to 1 to 3 lb. of white finishing gum to 5 gal. water.

A mixture that may be recommended for producing a soft, lustrous finish, and particularly for rayon braids, is given below:

a. Gum Arabic dissolved in 1
gal. Water 1 lb.
b. Gum Tragacanth dissolved
in 1 gal. Water 1/4 lb.

Use one part solution a and one part solution b to 4 to 5 parts water and apply lukewarm.

Running the goods through plain, lukewarm water and then through the calender will often remove wrinkles that have been developed in the process of dyeing.

Glue is the substance most often employed for stiffening braids, as well as other textile fabrics. This ingredient comes in many different qualities, and the grade required will depend upon the quality of the material to be treated and the result desired. The flakes or granules of glue should be allowed to dissolve in water some time before it is to be needed at the finishing machine, and as glue varies greatly, it is advisable to experiment with each new lot before proceeding with any quantity of material.

Various substances are used to prevent the size bath from souring. Among these are zine chloride, sodium fluoride, bluestone, and formaldehyde. Any of these chemicals are used in very small quanti-

Textile Size

Glucose			7	lb.
Soluble Oil			3	lb.
Magnesium	Sulphate		1	lb.

Textile Paste or Size

Potato Starch	100 lb.	
Calcium Chloride	300 lb.	
Water	300 lb.	

Manipulation: Soak starch and calcium chloride in the cold water for 2 hours then gradually heat mixture to boiling. Boil for 1 or 2 hours until a thick paste is formed.

Equipment: Clean wooden vat with open steam for boiling.

Cleaning Solvents for Textiles Formula No. 1

Carbon Tetrachloride

No. 2	
Carbon Tetrachloride	850 cc.
Heavy Benzoline, Purified	150 cc.

No. 3		
Heavy Benzoline, Purified	640	ec.
Ethyl Ether	120	cc.
Turpentine Oil, Purified	120	cc.
Ethyl Acetate	120	ec.
(Inflammable!)		
No. 4		
Heavy Benzoline, Purified	600	cc.
Turpentine Oil, Purified	120	cc.
Ethyl Ether	160	cc.
Ethyl Acetate	120	cc.
(Inflammable!)		
No. 5		
Carbon Tetrachloride	650	cc.
Alcohol	100	cc.
Ethyl Ether	100	cc.
Heavy Benzoline, Purified	80	cc.
Soap Spirit	50	cc.
No. 6		
Trichloroethylene		

Scouring Rayon Circular Knit Fabric

- 1. Run water in kettle (80-120° F.) using minimum amount that will enable the fabric to run freely over the reels. A properly loaded kettle of the correct type requires approximately a 20 to 1 bath.
 - 2. Load kettle with fabric.
- 3. Add 2 lb. soda ash or trisodium phosphate (depending upon water conditions).
- 4. Turn on steam and run goods for 10 minutes.
- 5. Add 3 lb. high grade neutral soap—olive or red oil base.
- 6. Add 2 lb. "soluble pine oil" or a similar solvent containing material. If desired this solvent material and soap can be added simultaneously in order to aid solvent dispersion.
- 7. Raise bath to boil. Observe condition of bath at all times. If bath does not show a good, clean, sudsy condition, add more soda soap and pine oil. It is impossible to accurately predict the amount of soda soap and solvent or the exact proportions of the same that will be required under an unknown set of conditions.
- 8. Run the kettle at or near the boil for 1 hour.
- 9. Drop bath and proceed with bleaching or dyeing operation.

Cleaning Tent Canvas

Mildew can be removed from a tent by sponging the canvas with a weak solution

of calcium hypochlorite, or bleaching powder. Be sure to wash the solution out well after using.

Cotton Textile Printing

For the shading of the pink print a paste is prepared with 40 parts of Irisamine G, that are dissolved in 400 parts of iron-free water. The resulting solution is then incorporated into 500 parts of starch tragacanth thickening, warming for a short time, agitating until the mass reaches 60-70° C, and entering 80 parts of acetate of chrome at 18° Bé, and bringing to 1000 parts through adding more water if this is necessary.

The starch tragacanth thickening, required in the above case, is prepared with 60 parts of wheat starch and 50 parts of wheat flour, that are made into a uniform semi-transparent paste with 700 parts of water, adding to this while still boiling 200 parts of a 6½% gum dragon muclage and 30 parts of olive oil. The bath being brought with water to 1000 parts

in all.

For the back of the pink print 200 parts of the above color paste are measured out and mixed first in a warm bath containing 800 parts of the starch tragacanth thickening, and then with 2 parts of acetate of chrome at 18° Bé. and 5 parts of acetic acid at 6° Bé., that are added at the right moment in the cooling

down bath. The shading product, needed for the red print, is obtained with 10 parts of a suitable brand of safranine, that are dissolved in 90 parts of acetic acid at 6° Bé., 10 parts of acetin and 300 parts of iron-free water. The resulting solution is then added into 500 parts of the starch tragacanth thickening indicated above, and after cooling sufficiently (60-70° C.) are entered 60 parts of a 50% tannin acetic acid solution and 40 parts of acetic acid at 6° Bé., bringing the whole to 1000 parts with further water.

For the backing of the red print a fourth printing paste is prepared with 2 parts of a suitable safranine, dissolved in 20 parts of acetic acid at 6° Bé, and 350 parts of iron-free water. The resulting solution is then poured into 550 parts of the starch and gum dragon thickening, and when this has been properly incorporated, steam is turned off, and the bath is left under the action of the agitator until 70° C. has been reached. Fourteen parts of a 50% tannin acetic acid solution, and 20 parts of acetic acid at 6° Bé, are then poured in, in close successions.

sion. The bath is made up after this to 1000 parts with further water.

The cotton cloth is printed with the above four color pastes, dried by passing through the hot-flue, and steamed for 1 hour without pressure, or for half this time with one half atmosphere. After the steaming, the goods are treated in a 1% tartar emetic bath at 50° C., rinsed for some time and dried. If the free acid in the goods is not eliminated in this way, the cotton cloth is passed through a second bath containing from 5 to 10 parts of chalk per 1. of iron-free water, giving a second rinsing, and drying and finishing.

If the printing is to be conducted on a pure white cotton cloth, the cost of treatment is much reduced, as a direct printing process is only required. This can be conducted with one of the pastes given below, the first of which requires, after its application and drying, a two hour steaming at 1 atmosphere pressure, while the second needs instead a one hour steaming with one-half atmosphere. Both colors being improved by a soaping.

Formula No. 1

Two and a half parts of alizarine black in paste S are mixed with ½ part of acetic acid at 6° Bé. and 6½ parts of a suitable starch thickening. The mixture is warmed until obtaining uniformity. After this it is allowed to cool down somewhat, and is entered ½ part of acetate of chrome at 20° Bé. bringing to 10 parts in all with water.

No. 2

Three hundred parts of a suitable brand of chrome orange are incorporated with 620 parts of acid thickening and 80 parts of acetate of chrome at 20° Bé., using the necessary precautions for avoiding loss of acetic acid. The acid thickening is prepared by boiling 210 parts of wheat starch with 570 parts of iron-free water, after having conducted properly the mixing in the cold. When a semitransparent adhesive has thus been produced steam is turned off, and toward 70° C. are entered 220 parts of acetic acid at 6° Bé., bringing with water to 1000 parts in all.

If the cotton material is colored in a light pink, this is obtained by dyeing on the jigger or on the padding machine with a suitable bath of Erika GN, shaded or not with Chrysophenine G; with a bath or Benzo fast scarlet 4BS (using the correct percentage), of Diamine rose BD, or of any other substantive pink; rinsing, drying and printing with the following

color paste:

Seventy-three parts of Ciba red G in paste are mixed with 27 parts of a 33% British gum thickening, and passed through a fine sieve, bringing then with water to 100 parts. Fifty-five parts of the above mixture are then entered in 12 parts of further 33% of British gum thickening, adding a little later 20 parts of caustic soda lye at 36° Bé., and 6 parts of glycerine. The whole is warmed just sufficiently for obtaining a uniform incorporation, and after having allowed the bath to cool down to about 50° C., are entered 71/2 parts of hydrosulphite NF concentrated, bringing with water to 100 parts.

When the cotton cloth goods have been printed with the above pink paste for obtaining the necessary details in the flowers and in the dark ground, the material is dried and steamed from 4 to 5 minutes at 105–107° C., being then left to hang for a short time, and finally treated with a bath furnished with 5 parts of olive oil or cottonseed oil soap and 2 parts of calcined carbonate of soda for every thousand parts of iron-free water, the bath being kept all through towards 60° C. After this the goods are given a last drying and are finished.

Logwood Speck Dye

Logwood Ex	tract	51°	Tw.	48	lb.
Soda Ash				30	lb.
Bluestone				12	lb.
This should	ho o	3;1,,+0	A + 5	about	0 30

Tw.

Seal Brown Cotton Dye

Cutch	35	lb.
Hypernic Extract	16	lb.
Logwood Extract	31/2	lb.

Add to dye bath and boil until dissolved, then add 3 lb. bluestone, add cold water, rake well and enter yarn. Give 6 turns and put down over night. Take up, give 6 turns, introduce into a solution of 4 lb. chrome at 160° F. and give 6 hours. Remove, wash well in cold water, put back in cutch liquor, 6 turns; into chrome, 4 turns; into cutch, 4 turns; into chrome, 4 turns. Wash off each time after chrome. Start new kettle with

Fustic Extract	7	lb.
Logwood Extract	31/2	lb.
Boil well for 2 hours.		

Violet Logwood Textile Ink
Logwood Extract (Weak) 300 lb.
Alum 12 lb.
Dextrin 15 lb.

Dissolve the alum by heating in a part of the extract solution. Finally 1½ lb. finely powdered lead acetate are slowly added and dissolved.

Textile Padding Liquor

Acetic Acid 50%	3	gal.
Formic Acid 85%	3	gal.
Glauber's Salt Crystals		Ĭb.
Water to make	100	gal.

The goods are padded on the face and the drying cylinders must not be too hot at first, so that sticking of the prints may not take place. Moderate drying in the initial stages should be the rule but at the same time if drying is not carried out properly there will be a grave danger of marking-off on the cylinders if any of the print color is allowed to adhere during the process. The wrapping of the first cylinder is sometimes advised in order to prevent sticking, but the circumstances in each case will dictate the precautions which will have to be taken. Two or three cylinders in any event will be found sufficient for the full development of the colors.

Preparation of Print Colors

In using the powder brands the following method of producing a print color is normally adopted.

Directoff Powder 8 16 cg

Dyesiun Powder	8-10	oz.
s pasted with		
Caustic Soda Solution		
(70° Tw.)	1/8-1/4	pt.
Monopol Oil or Similar		
Soluble Oil	1/8	pt.

Neutral Chromate Solution 1/4 pt.

The mixture is then allowed to stand for a short time before being added to

	 -		-
Water		2-3	pt.
and			-

Starch-Tragacanth 4-5 pt. (Thickening as Required) Making the whole up to

Printing Paste 1 gal.

For the production of lighter shades from the above standard a thickening of

the following type is made up.

Neutral Starch-Tragacanth 1 gal.

Caustic Soda Solution
(70° Tw.)

1 gal.

(70° Tw.) $\frac{1}{16}$ pt. Neutral Chromate Solution $\frac{1}{4}$ pt.

The neutral chromate solution is prepared in the following manner: Sodium Bi-Chromate

Crystals 1½ lb.

	TE	XTIL
dissolved in		
Water	6	pt.
To this add		_
Caustic Soda Solution (70° Tw.)	22	oz.
Make up to		
Neutral Chromate Solution		gal.
The paste brand dyestuffs as follows:	are pr	epared
Dyestuff	1	pt.
Neutral Chromate Solution	8,	oz.
Monopol Oil	21/4	pt.
Water Neutral Starch-Tragacanth	5	pt. pt.
Printing Color	1	gal.
		J
Thickening for Hand Print	ing or	a Silk
Formula No. 1		
Mix White Starch	5	lb.
and		
White Dextrin	5	lb.
with		
Acetic Acid, 12° Tw.	$7\frac{1}{2}$	
Olive Oil	2	lb.
and then add		
Water	$2\frac{1}{2}$	gal.
Boil to a paste.		
No. 2		
Mix White Starch	5	lb.
with	J	10.
Water	1	gal.
and		
Glue	$2\frac{1}{2}$	lb.
previously dissolved in		

Coloring Bone Articles

Boil to a paste, cool and add

Acetic Acid, 7° Tw.

21/2 gal.

lb.

lb.

Water

Olive Oil

Stir well.

The chief difficulty encountered in coloring bone material such as chess and other game counters, buttons, horn handles for umbrellas and walking sticks, ornamental vases and similar bric-a-brac of this type, etc., consists in obtaining good penetration of the dye. It is an unfortunate fact that certain acid and basic dyes of poor fastness to light will penetrate bone material better than some of the faster colors. Where penetration is too shallow, bone articles subjected to much handling like chess and draughtsmen, umbrella and walking stick handles and so on, soon disclose unsightly light

places where the superficial film of coloring matter has worn off.

Bone material is commonly dyed in a nested copper kettle, the inner container which carries the stock being perforated with small holes for the circulation of the liquor. The container can be lifted from the outer casing when it is desired to examine the stock during processing. Coloring of bone material is usually performed before it is polished, as treatment in hot liquor would roughen the surface of polished goods. When small articles like buttons, electric bell and light switch press-plungers, ivory sectors for inlay and marqueterie designs and so forth are to be colored, handling of the stock is facilitated by processing it in bags of linen net, each bag having a capacity of about 8 oz. of stock.

It is customary to boil-off bone material in clean water before coloring it. If the stock contains traces of oil or grease acquired during turning and fret-cutting of ornamental pieces, a small amount of pearl ash is put into the boil-off bath in order to emulsify the fatty substance. It is well to be sparing in the use of the alkali because the employment of an excessive amount will turn the bone a yellowish color. The use of soap for boiling-off is also apt to bring about this yellow discoloration in the stock; moreover, the presence of residual soap during coloring of the bone material will hinder penetration. The usual duration of the boil-off is from 15 to 60 minutes, according to the size of the pieces in the stock and the kind of bone. Antler and tusk material is harder and less porous than stock manufactured from sawn bone of bovine origin.

When the stock has been taken out of the boil-off kettle, it is plunged into the boiling dyebath, which is already fully charged with the appropriate dyestuff. Boiling proceeds for 30 to 60 minutes and then the stock is allowed to steep in the cooling bath for several hours in order to encourage penetration. It is not always advisable to process thin pieces made of horn at boiling temperature for longer than a few minutes, because of the risk of distorted material through softening of the structure in the hot liquor.

The following dyes may be employed for processing fast-to-light colors on bone material. Afterchrome Black of the PV type; Alizarine Brilliant Green G; Cloth Fast Yellow R; Eriochrome Red G; Erio Fast Brilliant Blue 3R; Radio Brown B; Cutch Extract; Logwood Extract. Afterchrome Black is applied to bone material in a boiling bath containing 1% of 30%

acetic acid. After processing for half an hour, 1% of sulphuric acid 168° Tw. is added and boiling is continued for a further half hour. The stock is then allowed to steep in the cooling bath for some hours, after which it is plunged into a fresh bath containing a boiling solution of bichromate of potash, the amount employed being from 1 to 2%. After 15 minutes processing at the boil, steam is turned off and the stock is left to steep for a further period of 15 minutes and then it is lifted and rinsed in warm water.

Alizarine Brilliant Green G yields fine blue-green hues of high fastness to light on clean white bone stock; when this color is used on discolored stock, or the darker sorts of horn material, the shade which ensues is a bottle-green color. Alizarine Brilliant Green G has good affinity for bone when applied in a boiling neutral bath. For deep shades with this dyestuff, an addition of 1% of acetic acid should be made to the bath after processing neutral for half an hour. Cloth Fast Yellow R also possesses good affinity for bone in neutral liquor. Deep hues may be processed with an addition of acetic acid, this to be put in when the bath has boiled for half an hour. Errochrome Red G yields rich red on bone stock. Dyeing should be commenced with the addition of 1% of acetic acid and when the bath has boiled for half an hour, 1 to 2% of bichrome may be put in. If the stock is hard tusk, boiling should be kept up for an hour before the bichrome is used. Erio Fast Brilliant Blue 3R produces a lively and very durable reddish-violet color on clean white bone material. This dyestuff has very good affinity for bone in a neutral bath. When processing a full shade, an addition of 1% of acetic acid may be made after the bath has boiled one hour.

Radio Brown B is a useful dyestuff for processing light or dark brown hues of first-rate fastness to light on bone stock. The affinity in a neutral bath is not good, hence an addition of acetic acid may be used at the commencement of dyeing. After the bath has been boiled for about half an hour, the color may be exhausted

by an addition of 1% of sulphuric acid. Cutch extract is an old favorite amongst bone dyes. This substance yields olive-gray to rich brown hues on bone, the shade depending on the processing method adopted. To produce olive-gray on bone stock, the material is boiled for 30 minutes in a bath containing 10 to 20% of dry cutch extract, and 1-2% of acetic acid. Steam is then cut off and

the stock allowed to feed in the cooling bath for 8 to 10 hours. The material is then put into a net bag and suspended in an empty barrel into which steam is blown for 10 minutes. The jet of steam must not impinge directly upon the stock. Oxidation of the cutch which has been absorbed is then completed by exposing the bone pieces to the air while they are spread out in shallow trays. In order to develop the olive-gray coloration, the stock is plunged into a boiling bath containing 2 to 5% of green copperas. Steam is cut off after the material has boiled for 15 minutes, after which the stock is left to steep for half an hour and then rinsed. The olive-gray hue produced in this manner has long been a popular color for the bone platings on pocket knives. If it is desired to process orangebrown or deep reddish brown with cutch, development of the color is done with bichrome and copper sulphate instead of green copperas. When deep colors are being processed on bone material with cutch, or other natural coloring matters, it is usually necessary to remove the film of loose color and resinous impurities which forms on the surface of the bone during processing. If this film is not cleaned off, it clogs in the bone and hinders development of the final color during after-treatment with the metallic salts. In order to cleanse the stock, the pieces are put in the loose condition into a tumbler apparatus containing a thin paste of sawdust and water, or preferably cow dung and water. When the device is set into motion, the movement of the stock in contact with the sawdust, etc., cleanses away the film.

Logwood extract is sometimes combined with cutch for the purpose of modifying the tone of the latter. Logwood extract is also used for deep black on bone articles, the process consisting in boiling the stock in a solution of logwood extract, followed by the oxidation of the hematine by steaming and exposure to the atmosphere. After the material has been freed from film in the tumbler apparatus, the black color is developed in a boiling bath containing copper suphate and green copperas. A black of this kind is not as fast to light as afterchrome black, but penetration is frequently better than in the other instance.

Dyeing Vegetable Ivory Buttons

The following is suggested with the use of basic dyes: The buttons are boiled in water for 1-2 hours before dyeing. Pale shades are dyed for 2 hours at the boil in

a neutral bath; if the water is very calcareous, some acetic acid must be added.

Full shades are first mordanted for 4 hours in a bath prepared with 40 parts tannin per 1000, then rinsed in cold water and treated in a bath prepared with 20 parts tartar emetic per 1000 for ½ hour at 120–140° F. The buttons are then rinsed for ½ hour with boiling water in order to remove the free mordant and dyed in a fresh bath acidified with acetic acid.

Dyeing Brush Bristles

When dyeing fiber materials to be used for the manufacture of brushes, etc., and necessitating the material being dyed through well, it is best to use a combination of about 2-3% of a direct black

and 2-4% logwood extract. Charge the starting bath with 2% ammonia and 1/4-1/2% soda ash, add 2-3% dye previously well dissolved in condensed then about 5% cryst. water, and Glauber's salt; boil up well, enter the material, work for 5-10 minutes, cover with a lattice frame weighted with stones, boil for 2-3 hours, and allow to feed for 1/2-1 hour in the cooling bath. Then lift the material, allow it to lie exposed to the air for several hours, and enter into a fresh bath heated to 30-40° C. (85-105° F.) containing pyrolignite of iron of 4-7° Tw.; leave in this bath for ½-1 hour, throw out and leave exposed to the air for several hours, rinse well and dry.

If so-called patent or luster-fiber is to be produced, the method of working is exactly as described above; only the fiber is finally taken through a bath of 40-50° C. (105-120° F.) charged as follows:

- (,			
Liquor		10	gal.	
Gelatin Glue		2	Ϊb.	
Soft Soap		2	lb.	
Logwood Extra	ct	2	lb.	
Fustic Extract		1/2	Ib.	
Pyrolignite of	Iron	1/5	lb.	

Treat the goods in this bath for 30 minutes, allow to drain, and brush dry with suitable brushing machines. If the fiber is not lustered, 8 oz. of whitening per 10 gal. liquor are added to the bath of pyrolignite of iron.

The dye liquors may be used repeatedly; dyeing in the standing bath requires about ½-2% of the stated quantities of dye and logwood extract, equal quantities of soda and ammonia, and about 3% salt calculated on the weight of the goods.

Coconut Fiber Dveing

Dyestuff	30	lb.	
Acetic Acid, 30%	90	lb.	
Glycerin (only where the			
goods will be steamed after			
printing)	30	lb.	
Water	400	lb.	
Thus one countly Inhistory in a	150	71.	

Tragacanth Thickening If the mats are to be steamed, the operation is carried out in a cottage steamer, the duration of steaming being from a quarter to half an hour without pressure. The mats are hung on rustless metal hooks riveted into movable metal strips which span the interior of the steaming cottage. The stock is seldom washed after steaming, unless the thickening has been made too good with the result that the printed portions handle stiffly. Basic dyes are apt to lose depth during washing, even when the stack has been steamed; hence, washing is only done where the necessities of the case call for it. Some printers regularly make an addition of tannic acetic acid to the print color in order to heighten the resistance of basic color to washing and to general wear in the domestic sphere. The following basic colors are suitable for use in printing coir matting: Phosphine, rhodamine, magenta, safranine, methylene blue, malachite green, methyl violet, bismarck brown, jute black.

Substantive dyes prove useful for printing coir in designs of good fastness to washing. This class of dyes should be steamed after printing in order to obtain good results. The printing paste is made as follows:

 Substantive Dyestuff
 30 lb.

 Water
 370 lb.

 Phosphate of Soda
 30 lb.

 Glycerin
 70 lb.

 Tragacanth Thickening
 500 lb.

 (40:1000)
 500 lb.

The following substantive colors are suitable for printing coir: Chrysophenine G, Direct Fast Scarlet 4BS, Benzopurpurine 4B, Direct Bordeaux 6BS, Direct Brown G, Direct Brown M, Direct Fast Pink BK, Direct Green B, Direct Sky Blue FF, Direct Black BH, R, E. After the mats have been printed, they are allowed to become partially dry and then they are steamed without pressure for half an hour. They are then rinsed in cold water.

Bleaching Coconut (Coir) Fiber

The bleaching process with hypochlorite is carried out in a cold bath after the coir stock has been boiled out in a solution of caustic soda. From 3 to 7 lb. of

commercial hypochlorite of soda solution are used per 100 gal. of water in the bleach bath. The stock is allowed to remain in the kettle for from 1 to 8 hours after which it is soured in a fresh, cold bath containing 11/2 pt. of hydrochloric acid, 30 to 34° Tw. per 100 gal. of water, and subsequently well rinsed. The batch is then ready for antichloring, this process consisting of immersing the coir for a period of 10 minutes in a fresh, cold bath charged with 11/4 lb. hyposulphite of soda crystals per 100 gal. water. After this has been done, the stock is thoroughly rinsed in cold water, then steeped for several hours in two or three changes of water and finally centrifuged.

To bleach coir stock with permanganate of potash and bisulphite of soda, the material is first boiled out in a kettle with 3% caustic soda and after being rinsed, it is immersed for 12 hours in a cold solution of permanganate of potash, 34° Tw. The stock is then rinsed and entered into a fresh cold bath containing a solution of bisulphite of soda 34° Tw. When the stock has steeped for one hour, the bath is let down, the material being then given two cold rinses. If it is then found that decolorization is insufficient, the operations just outlined are repeated.

In a case where hydrosulphite is chosen as the decolorizing agent, the stock is first soaked in cold water for 24 hours to remove the looser class of impurities and then a liquor containing 10 to 15 lb. of hydrosulphite per 100 gal. of water is prepared in a separate kettle connected by piping to the other one. The solution of hydrosulphite is then run in at a temperature of about 85° F., circulation of the liquor being kept up for 20 minutes or so by means of a rotary pump attached to the apparatus. After this period has elapsed, steam is turned on and the kettle is raised to about 170° F. and maintained at this temperature for from 1 to 4 hours. If the stock is heavily colored with natural pigment, further amounts of hydrosulphite are added to the kettle from time to time. When decolorization is deemed sufficient, the bath is let down and the stock is well rinsed in cold water.

Some manufacturers of coir mats prefer to decolorize the stock in the woven condition. In this event, the mats are either strung on rods which rest upon the rim of the kettle or else they are processed in a package apparatus. This is of an extremely simple type, it consisting of little more than an open kettle fitted with a rotary pump for circulation purposes. It is customary to place a wooden trammel or grid on top of the

pack to circumvent floating of the stock due to the formation of steam pockets.

Bleaching Vegetable Fibers German Patent 615,680

Steep for 10 minutes in hot water and then place in bath containing 2.2 g. active chlorine and 1.5 g. caustic soda per l. at 32° C. Raise temperature to 75° C. and treat with hydrogen peroxide, then rinse.

Bleaching Mohair Cotton Fabric

The cloth, which is first thoroughly scoured in a soap soda ash bath, is transferred to a winch containing 500 gal, of water at 100° F. Five lb. of potassium permanganate carefully dissolved in lukewarm water are slowly added through a fine sieve. The cloth is run in this bath for 11/2 hours. After two cold 10-minute rinses the box is filled to the same height as before with cold water and 4 gal. of 72° Tw. sodium bisulphite liquor are added. The cloth is run several minutes before adding 12 lb. of commercial sulphuric acid previously diluted by pouring into several times its volume of cold water. The cloth is run in this bath for 2 hours. A wash in a bath made slightly alkaline by adding trisodium phosphate, followed by a thorough rinse completes the process. It is sometimes necessaary to add a small amount of Acid Violet, Color Index No. 698, to the last rinse to obtain the bluish white which is usually requested.

Potassium permanaganate also has a limited use in producing novelty effects on shoe plush. The shoe plush after a good scour is dyed brown by running in a bath containing 30 lb. of permanganate per 825 gal. of water at 120° F. for 11/2 to 2 hours. An addition of 5 to 10 lb. of potassium permanganate is usually necessary to obtain the desired depth of shade. Following the dyeing the cloth is rinsed at 160° F. with water made slightly alkaline by adding 1½ lb. of trisodium phosphate. Two warm rinses complete this part of the process. The novelty two-colored effect is obtained by using a brush tipping machine. The latter is essentially a one-color printing machine which uses a brush roller instead of an engraved roller. The pile is tipped with an acidulated solution of hydrogen peroxide. If nothing more is added to the tipping liquor a brown pile with a lustrous white tip is obtained. By adding certain basic and acid colors not affected by the peroxide beautiful blue, green and rose tips over a brown base are obtained.

A gray, varying in intensity from a light rabbit's fur color to a jet black, can be substituted for the brown at the base of the pile. The depth of the gray is directly proportional to the depth of the manganese brown originally on the fiber. It is accomplished by immersing the cloth after the tipping treatment in a cold bath containing .5 to 12.5% aniline salt and .25 to 12.5% sulphuric acid, depending on the depth of shade desired. It is worked in this bath for 30 minutes. A weak ammonia rinse and a thorough wash completes this process.

The above principle—aniline black over manganese brown—is sometimes utilized to obtain clear white discharges on

woolen fabrics.

Eleaching Yarns, Skins and Straw U. S. Patent 1,966,915

One hundred grams of woolen yarn may be placed in a solution of 1000 cc. of methyl alcohol in which 30 cc. of hydrogen peroxide (30% water solution), are incorporated. As the oxygen of the hydrogen peroxide is liberated much more freely in an alkaline solution, there should also be added about 2 cc. of, preferably, concentrated ammonia water. The solution containing the yarn should be heated to about 60° C. for about 8 hours.

The pelt is put into a bleaching liquor of about 1000 cc. of ethyl alcohol containing about 15 cc. of hydrogen peroxide (30% water solution), about 0.3 cc. of concentrated ammonia water, and about 45 g. of Turkey red oil. The skin is allowed to remain in the bleaching liquor for 24 hours at about 18° C. The skin thus treated exhibits perfect bleaching and the complete absence of injuries or impairments.

Pandan "stumps" are treated with a 1000 cc. ethyl alcohol solution containing 35 cc. of hydrogen peroxide (30% water solution) for about 6 hours, at about 60° C., and are then finished in the usual way. The bleaching proceeds very smoothly because the chlorophyl is ex-

tracted by the alcohol.

Natural	Finish	for	Calico	
---------	--------	-----	--------	--

Potato Starch	5 lb.
Wheat Flour	7½ lb.
are boiled with	
Water	250 lb.
then add	
China Clay Paste	10 lb.
and	

French Mineral White	10	lb.	
Boil and add			
Coconut Oil	3/4		
White Soap	1/2	lb.	
Carbonate of Soda	1/4	Ib.	
Water	3 1	lb.	
Add to a vat containing			
Potato Starch	15	lb.	
and			
Water	75	lb.	
Stir thoroughly and then s	slowly a	add	
Potato Starch	5	lb.	
and			
Water	5	lb.	
with a trace of wilthamanine			

with a trace of ultramarine.

The starched goods are dried in a dry room, damped and rolled under pressure.

Alizarine Lake Formula No. 1

Sulphate of Alumina (Tech. 18% Al_2O_3) 972 lb. Water 10,000 lb. No. 2

 Soda Ash
 500 lb.

 Water
 5,000 lb.

Filter both solutions.

Add the hot soda solution slowly to the hot alumina solution while stirring, keep boiling gently until the precipitate begins to be glassy, wash with clean water free from iron until, by repeatedly decanting, a sample of the wash water shows but very little turbidness with chloride of barium solution. The alumina now obtained by filtering may be used at once for making alizarine lake. The weight of the paste filtered into the bag amounts to about 7000 parts. Add to the alumina paste a solution of 144 parts calcium chloride anhydrous, chemically pure, in 500 parts water, and follow, while stirring well, with a solution of 84 parts phosphate of ammonia (pure neutral salt) in 500 parts water. Then stir in 150 parts ammonia Turkey red oil, which has been previously dissolved in a little water, and finally add 1000 parts Alizarine Red 1B extra (20% paste).

Either boil this preparation for 6-10 hours in an open vessel, when the evaporated water must be replenished, or treat for 1 hour in the autoclave with about

59 lb. pressure.

Alizarine Cyclamine is affected by metals including copper, and for this reason should not be steamed in the autoclave; lead vessels, however, may be used without risk.

Every substance used in the making of

madder lakes, including the water, must be free from iron.

Alizarine Dyeing of Silk

a. The well cleaned silk is entered, worked and steeped over night in a cold bath of basic aluminum sulphate prepared by dissolving 171/2 oz. aluminum sulphate, free from iron, in 1 gal. of water to which 4 oz. soda crystals dissolved in a pint of water is added, the clear solution showing 12-15° Tw. The silk is wrung out from the mordanting bath, rinsed well, then fixed for half an hour in a cold bath of sodium silicate of 1° Tw. and finally rinsed very thoroughly. A basic aluminum salt is thus obtained on the fiber without injuring any of the properties of silk. The mordanted silk is then dyed with alizarine paste, the quantity of alizarine used depending upon the depth of the shade to be dyed, in a boiled off liquor bath broken with acetic acid, entering and working it in the cold for half an hour, gradually raising it to the boil in 1 hour and dyeing at that temperature for another half an hour. The dyed silk is then thoroughly washed in water, brightened in a weak bath of acetic acid and finally dried. The silk is dyed bright red.

b. Silk after being properly cleaned is entered, worked and steeped overnight in a cold bath of "nitrate of iron"—basic ferric sulphate—32° Tw. It is wrung out the next morning from the mordant bath, rinsed well in water, then fixed by working for half an hour in a cold bath of sodium silicate of 1° Tw. and finally rinsed very thoroughly in water. This mordanted silk is dyed with alizarine as usual. This gives a bright violet color.

c. As chrome cannot be used with advantage on silk as with wool, on account of its tendency to destroy the luster and injure the fiber, the mordanting is usually done with chromium chloride or chromium sulphate. The well scoured silk is worked and steeped overnight in a cold bath of basic chromium chloride 32° Tw. The next day the excess liquor is squeezed out, the mordanted silk is well washed in water, fixed for half an hour in a cold bath of sodium silicate 1° Tw. and finally rinsed very thoroughly. A basic chromium salt is thus obtained as a mordant on the fiber without particular injury to any of the properties of silk. The mordanted silk is then dyed in a boiled off liquor bath broken with acetic acid as The dyed silk is thoroughly washed, brightened with acetic acid and dried. Silk is dyed a bright chocolate color.

Chrome Dyeing Formula No. 1

Chromium Black	7	ĸg.
Acetic Acid	4	kg.
Heat to 100° C., boil ½ hour,	add:	_
Formic Acid	1	kg.

boil another ½ hour, add
Potassium Bichromate

at a temperature of 70° C., then go up to 100° C., and boil ½ hour.

1.5 kg.

2 kg.

No. 2

Chromium Blue	4	kg.
Acetic Acid	5	kg.
Glauber's Salt	10	kg.
Heat to 100° C., boil 1/2 hour,	add	
Formic Acid	2	kg.
		-

boil ½ hour more, add

Potassium Bichromate

1.5 kg.
at 70° C. up to 100° C., boil ½ hour.

No. 3

Chrome Flavin 2 kg. Additions, method as in No. 2.

Vat Dyeing

rormula No. 1	
Vat Scarlet	4 kg
HN-Process, 50-55° C.	Ŭ
"One bath" process.	
Fundamental Vat	
Dyestuff	1 kg.
737-40	20 1

Water	30 kg.
Caustic Soda (40° Bé.)	1 kg.
Hydrosulphite	1 kg.
Dyeing Vat	
Glue	3 kg.
Ammonia	3 kg.

Hydrosulphite
Dye ½ hour at 50-55° C.

No. 2

Vat Scarlet 6 kg. HN-Process, 50-55° C. "Two bath" process.

Fundamental Vat As in No. 1.

Dyeing Vat As in No. 1.

Dye ½ hour at 50-55° C., then add 2% hydrosulphite for the second (application), dye another ½ hour at 50-55° C.

	TEXTILE
No. 3 Vat Black HN-Process at 50-55° C. "One bath" process.	2 kg.
Dissolve the solid vat (kii the same amount of boiling wa 5% (of the dyestuff weight the same of hydrosulphite.	ater, adding
Dyeing Vat Glue Ammonia Hydrosulphite Dye ½ hour at 50-55° C.	3 kg. 3 kg. 2 kg.
No. 4 Vat Black HN-Process, 50-55° C. "Two bath" process.	12 kg.
Solid vat solution (see No	3).
Glue Ammonium Sulphate Hydrosulphite	3 kg. 4 kg. 2 kg.
Dye 1/2 hour, then add: Hydrosulphite Ammonium Sulphate Repeat dyeing 1/2 hour (se	2 kg. 3 kg.
No. 5	
Vat Blue HW-Process, 60-65° C. "One bath" process.	3 kg.

Dyeing Formula for Acetate Rayon Velvet

2.2 kg.

1

30

1

kg.

kg.

kg.

kg.

kg.

kg.

Fundamental Vat (Stammküpe)

Caustic Soda (40° Bé.)

Dyestuff

Hydrosulphite

Hydrosulphite

Dye 1/2 hour.

Water

Dyeing Vat Glue

Ammonia

Formula No. 1

Substantive Dyestuff	2 lb.
Glycerin, Dynamite	4 lb.
Condensed Water	2 gal.
British Gum Thickening	6 gal.
Caustic Soda, 75° Tw.	1 gal.
The following is an example	of a print
color for acetate rayon velvet.	
m	7 77.

Basic Color 1 lb. Acetic Acid (30%) 20 lb. British Gum or Senegal Thickening 8 gal.

A proportion of tannic-acetic acid, 1:1 improves the fastness to washing, in deep shades

The following is an example of a formula for a print color containing tannic acid:

Basic Color	2	lb.
Acetic Acid, 30%	15	1b.
Acetine	$1\frac{1}{2}$	lb.
Water	5	gal.
British Gum	18	lb.
Tannic-Acetic Acid 1:1	10	lb.

The last named ingredient should be added only when the color has become cold.

After steaming, the pieces are treated for a few minutes in a lukewarm bath charged with 12 oz. of tartar emetic per 10 gal. of water. This operation is commonly performed in a star machine, but it may also be carried out in a winch apparatus where the more robust velvets are being handled. After being treated with tartar emetic, the batch is given a light rinse in cold water, after which the pieces are hydro-extracted.

Vat Printing Color

Paste Vat Color	10	lb.
Glycerin, Dynamite	$3\frac{1}{2}$	lb.
Carbonate of Potash	14	lb.
Sodium Formaldehyde Sul-		
phoxylate	7	lb.
British Gum Thickening	7	oral.

The following recipe for a color for the brush printing of viscose rayon plush will furnish an indication of the proportions of substantive dyestuff and other ingredients used in preparing the print colors:

Diphenyl Brown BBN Extr	a 8	oz.
Direct Orange G	3	OZ.
Chrysophenine G	8	oz.
British Gum (Dry)		oz.
Glycerin	10	oz.
Phosphate of Soda	12	oz.
Condensed Water	1	gal.

Wool Dyeing

Indigo (20% Paste)	10 lb.
Water	2.4 gal.
Sodium Hydrosulphite	
(Powder)	2.5 lb.
Caustic Soda (76° Tw.)	6 pt.

The indigo and the water are intermingled first. To this mixture, the sodium hydrosulphite is added, gradually and with unceasing stirring. Finally, the caustic soda is introduced. The mixture is to be frequently stirred and its temperature maintained at 60° C. In about two hours, complete reduction may be expected.

Indigo Fermentation Vat Formula No. 1

Indigo (60%)		20	40	lb.
Woad		560 - 1	120	lb.
Bran or Sharps		30	40	lb.
Madder		10-	15	lb.
Lime		12-	25	lb.
Water		3	240	gal.
No	. 2			_
Water		21	60 g	gal.

No. 2	
Water	2160 gal.
Woad	5 cwts.
Natural Indigo (Paste)	20–40 lb.
Bran	5 buckets
Madder	6 lb.
Lime (In Slaked Form)	3 gal.
Lime (In Slaked Form)	as directed

The water is run into the vat and raised to the temperature of 135° F. The woad is now added and the liquor stirred several times till "pasted." The 3 gal. of slaked lime are stirred in and he whole left over night.

A representative British hydrosulphite vat for wool may be made up in accordance with the following tabulation:

Water	1080 gal.
Ammonia (25%)	3.6 pt.
Hydrosulphite Powder	$2 ext{ lb.}$
Glue Solution (1:10)	2.4 gal.
Indigo Solution (20%)	2.4 gal.

The water is run into the vat and the temperature brought up to 120° F. The indigo solution is stirred in.

At the beginning of each dyeing operation, add ammonia, hydrosulphite powder and indigo solution. At the end of the day's run, add a little glue solution and 1.2 qt. of caustic soda (at 76° Tw.).

Printing of Animal Fibers U. S. Patent 1,962,601

Colored patterns fast to washing, light, perspiration, etc., are obtained by printing prechlorinated wool and silk with a thickened paste containing Indigosol, Leucosol, or similar water solvent derivatives of vat dyes and sodium nitrite, then steaming with wet steam at 99-100° C. for 7 minutes, and passing the fabric in open width through dilute sulphuric acid (50 g. [density 1.53] per l.) at 95°, followed by washing and oxidation with a solution at 35-40° containing (per l.) 1.5 g. of sodium persulphate and 2 g. of sulphuric acid (density 1.015) for 20 minutes.

Dyeing Aged Black on Piece Goods
The following is suggested: 120 lb.
aniline salt, 10 lb. aniline oil, 35 lb.

sodium chlorate, ½ lb. copper sulphate, per 100 gal. liquor.

The goods are impregnated with this solution, aged and chromed.

The following is another method of dyeing an "ungreenable" aged black. Two solutions are prepared:

a. 55 gal. of water, 45 lb. aniline salt,
13½ lb. toluidine, 7 lb. acetic acid, 18½
lb. sodium chlorate.

b. 18½ lb. nitrate of iron, 76.6° Tw., 6 gal. water, 27 lb. of a solution of copper sulphate (2:10).

Mix 8 gal. of a with 1 gal. of b, and pad with this mixture. Age and develop as usual.

The following process is also recommended: The pieces are padded with the following solutions, which are prepared separately, mixed when cold, and made up with water to 100 gal. The padding liquor should stand at 12° Tw.; 120 lb. aniline salt are dissolved in 26 gal., 3½ pt. water; 5½ lb. copper sulphate are dissolved in 10 gal. water; 37 lb. 9½ oz. sodium chlorate are dissolved in 7 gal. 3½ pt. water; 4 lb. ammonium chloride are dissolved in 2 gal., 3½ pt. water; to this are added 4 gal., 6½ pt. aluminum acetate, 15° Tw.

The cloth should be impregnated in such a manner that it retains about its

own weight of padding liquor.
After impregnation, the cloth should be dried as rapidly as possible at a low temperature, after which it is aged for 1 to 2 hours at a temperature of 92° to 96° F.

The aging is followed by chroming and soaping.

Cotton Printing Paste

Victoria Blue B		6	oz.
Methyl Violet 4B	•	$\frac{1}{2}$	oz.
dissolve in			

Acetic Acid, 40% ½ gal.
Starch Thickening (1 lb.
Wheat Starch/1 gal.) 5 gal.
when cold add

Tannic Acid (4 lb./1gal.) 1/2 gal.

Crimping Cotton

Beautiful effects may be obtained by printing on a Gum Resist and subsequently passing the cloth through strong caustic soda. The dry content of the gum used as a resist is very important. A very highly converted British Gum is usually used and the strength will run 3-4 lb. per gal.

The greater the dry content of a gum

resist, the more effective is its power to resist the caustic soda. The latter will vary in strength from 25 to 30% according to the length of time the cloth is let lie after immersing and squeezing and prior to washing out. For best results it is advisable to select a light weight cotton cloth and print a design that is largely composed of lines running parallel to the selvage of the cloth. The reason for this is that the shrinkage, for the most part, takes place in the warp. After printing, run the cloth through the strong caustic soda in a pad box and let set 1 to 2 minutes. Finally rinse well with cold and hot water, hydro-extract and dry in a crepe dryer. In dyeing grounds for this type of work it is best to select colors that will not be affected by the caustic scda. If crepe dyeing is possible then beautiful two-toned effects may be obtained by dyeing the cloth after crimp-

In dyeing the latter, the dyestuff will have much more affinity for that part of the cloth that has been attacked by the caustic and as a result this portion will come out much heavier. Other effects may be obtained by selecting printing colors that will develop in a steaming operation and that will work well with the Gum Resist. These colors are printed on with the Gum Resist and then steamed, padded with the caustic and finished as mentioned above. The final result is a crimp in the colored or printed portion of the cloth. By selecting dyed grounds that may be discharged it is possible to obtain a crinkle in the white portion of the cloth. A discharge is made up with the Gum Resist and upon printing and steaming, the color is discharged at the printed part. After running through the caustic soda and finishing as mentioned above, it will be noted that the crimp is in the white portion of the cloth whereas the colored portion is uncrimped.

Lacquer Printing of Cloth with Metallic and Pigment Colors

This type of work is largely being carried out on silk, rayon and celanese where excessive handling is to be avoided. The advantage of this type of printing is in the fact that finished goods may be printed, dried and shipped without any intermediate process of steaming, washing, etc. The colors are really in a sense painted on the cloth and the secret of the success of this type of printing lies chiefly in the softness of the resultant print. Formerly bronze and pigment prints were extremely harsh when printed

by this method but today the lacquers used have been highly developed and the prints are much softer in feel. Both cellulose acetate and nitrocellulose lacquers are used and the difference between the two is very slight as far as the resultant print is concerned.

Bronze or metallic prints are nowheres near as fast as the pigment class of colors. They tend to go dull on standing and will wash out in time. Pigment colors are extremely fast and will even stand a good rubbing. In order to do a perfect job, the engraver, printer and colorist must work together. The engraving is very important as too shallow a depth will make the color stick-in. The colorist must have the proper amounts of solvents in his printing paste, so that the paste will not dry too fast in the engraving. The printer must run at a uniform speed so that the paste as worked out by the colorist will give even results. Too fast a drying paste will make the color stick in, whereas too slow a drying paste will not dry fast enough over the dry cans. A nitrocellulose lacquer can be made by dissolving the dry nitrocellulose in a mixture of acetone and ethyl or methyl acetate. A cellulose acetate lacquer can be made by dissolving the dry substance in a mixture of alcohol, phenol and solvent naphtha. In using pigment pastes it is advisable to have them extremely finely ground in some solvent, such as acetone together with olive or castor. Proper grinding requires special equipment and this treatment is very essential for the best results.

	Aniline Black Printing I	aste	
	Yellow Prussiate of Potash	8	oz.
	Chlorate of Soda Crystals	5	OZ.
	dissolved in		
	Hot Water	2	pt.
	and added to		
-	Aniline Salt	12	oz.
	previously dissolved in		
	Hot Water	1	pt.
	and stirred into		
	Starch Tragacanth Thickenin	ng	
	Printing Paste	5	pt.

Silk Printing Pastes Formula No. 1

Five pounds of good white starch and 5 lb. of white dextrin are mixed with 1 gal. of water, 7½ lb. of acetic acid of 12° Tw., 2 lb. of olive oil and 2½ gal. of water are then added, and the whole

boiled into a paste. This will suit almost all colors.

No. 2

Five pounds of good white starch are mixed with 1 gal. of water, and 2½ lb. of pale glue previously dissolved in 2½ gal. of water are added, and the whole boiled up to a paste. After allowing to cool, add 5 lb. of acetic acid 7° Tw., and 2 lb. of olive oil.

Textile Printing Pastes Formula No. 1

Wheat Starch Thickening

Wheat Starch 12 oz. Water 6 pt.

Boil and add

Chlorate of Potash 6½ oz.

When dissolved add

Yellow Prussiate of
Potash 8 oz.
Aniline Salt 12 oz.
Aniline Oil ½ tumbler
Printing Paste 1 gal.

No. 2

Copper Sulphate Black for Block Prints (Thickening)

Water 4 gal.
Boil together and when thickened add
Gum Tragacanth 1 gal.
Boil further until an even texture is pro-

duced, cool and make up to
Printing Paste 8 gal.

Use 7 parts of thickening to 1 of aniline hydrochloride.

Silk Printing Color Resist

Rosin		650 lb.
Yellow Wax		50 lb.
Spermaceti		30 lb.
Suet		18 lb.
Paraffin		25 lb.
	-	

Turpentine Oil 250 lb.

are heated together until they form a thoroughly liquid mass.

This resist is printed on lukewarm either in the printing machine with very deeply engraved rollers, or by hand printing. For the latter purpose the above mass must be kept a little thinner by the addition of a little more turpentine oil.

After printing, the goods are sprinkled with fuller's earth, and then hung up for a few days at the ordinary temperature.

When the resist is dry, the goods are washed in cold water and dyed in a cold bath.

Fancy Textile Printing "Resists"

The following is a good and simple formula generally used by textile printers. It washes with water.

Formula No. 1

English Drop Black or

Lamp Black ¼ oz. Paraffin Oil

enough to make into a paste

Amalgamate the above thoroughly and add:

Tincture of Green Soap 4 oz. mix and add:

Concentrated Lye Solution 5 drops Stir vigorously and keep in well corked bottles.

No. 2

Japan Color
Raw Linseed Oil
Boiled Linseed Oil
Washes off with oil solvents.

No. 3

Powdered Castile Soap 1 oz. Hot Water 2 oz. dissolve thoroughly and add show card color.

White Resist for Sulphur Dyes

Write Resist for Sulphur Byes
British Gum
200 g.
Water
250 g.
Zinc Chloride
400 g.
Water
150 g.

Heat until the gum and salt are dissolved.

If the resist white is found to run when printed with heavily engraved rollers, it may be improved by the addition of 75 to 100 g. China clay per kg. of color: as a rule, however, this addition is not necessary with ordinary patterns.

Cotton Yarn Dye Resist

If cotton is first impregnated with a solution of tannic acid overnight, squeezed in the morning, and then immersed in a bath of stannic chloride, it takes on the property of resisting many dyes. Use about 3 lb. of tannic acid (on 100 lb. of cotton yarn), and fix in a bath containing 2½ lb. of stannic chloride crystals. This tannic acid bath is used hot at the time the cotton is entered, but is cold by morning. The tin bath is used cold.

Wax Resist for Woolen Yarn

Rosin	60 lb.
Yellow Beeswax	5 lb.
Mutton Suet	2 lb.
Spermaceti	3 lb.
Paraffin Wax	2 lb.
Turpentine	4 lb.

The above are heated toge her and the resulting paste is printed on the goods. Strew with fuller's earth to prevent sticking, and when dry, wet out the skeins in cold water and dye in a cold or lukewarm bath with the required acid color.

Acid and Alkaline Resistant Treatment for Wool

U. S. Patent 1,964,934

Sulphite Cellulose Waste		
Liquor (Lime Free)	90	oz.
Magnesium Chloride	10	oz.

Stripping Sulphur Colors from Mixed Fabrics

A simple and yet very effective way of stripping sulphur colors on cotton in the presence of wool or worsted is as follows:

Prepare a cold bath of 1½° Tw. chloride of lime. Run the cloth full width in this bath for 30 minutes. Then drop the bath and rinse thoroughly with cold water. A second bath containing ½° Tw. commercial hydrochloric acid is now made. The cloth is run in this bath for 20 minutes at 160° F. and then rinsed thoroughly. The excess acid is finally neutralized by a run in a lukewarm ½% soda ash solution. A final light soap scour completes the process.

The chemic treatment should destroy practically all the sulphur dyestuff inside of 15 minutes if the chemic is freshly made. When old chemic solutions are used longer running or a stronger bath is necessary. The hydrochloric acid treatment removes any residue of rust or sulphide spots left from the chemic treatment. The resulting cloth is usually a light cream color which the soap scour makes considerably lighter. The wool is chlorinated slightly by this method and has an increased luster and a greater affinity for dyestuffs.

Hydrosulphite Discharge on Indigo Ground

The printing paste is prepared as follows:

Hydrosulphite NF Concentrated 125-200 lb.

are stirred into

Hot British Gum
Thickening 655-580 lb.
and after cooling

Zinc White Paste 1:1 150 lb.

Anthraquinone Paste 30% 50 lb.

Acetine (Neutralized with Soda) 20 lb.

are added.

The amount of hydrosulphite in the discharge depends upon the depth of the indigo shade.

After printing, the goods are well dried and then steamed for 3 minutes at 216–218° F. in the Mather-Platt, which must be free from air. The washing of the steamed goods is best carried out at full width in the washing machine in a boiling bath containing 10 parts silicate of soda 66° Tw. to 1000 parts water and 3 parts formaldehyde 40%. The passage through the washing machine should take three-fourths to one and a half minutes and the goods then well rinsed.

Instead of washing with silicate of soda, quick-lime (5 parts per 1000) or caustic soda solution may be used, although the silicate has the least effect

on the indigo bottom.

It is advisable to steam and finish the printed goods as quickly as possible, but if this cannot be done immediately, the material must be protected not only before but also after steaming against moist air by winding rolls and keeping in a warm dry room 85-100° F. After steaming the white is cleared as above by passing the pieces through an alkaline bath.

Although the indigo is readily converted into a leuen-compound by hydrosulphite, still the discharged places are apt to show a bluish tint if the reduced compound is not completely removed from the printed parts, or if the indigo-white is partly reoxidized to blue before the steamed pieces are washed. The addition of anthraquinone to the printing paste aids the discharging effect of hydrosulphite and prevents the indigo-white from being too quickly reoxidized.

Crease Proof Fabric British Patent 424,535

Ammonium sulphocyanide in the presence of variable quantities of urea has the advantage of requiring a comparatively low temperature for its formation. In previous similar processes it has been found necessary to heat the resin mixture for several minutes at 160

to 180° C. in order to produce full polymerization, but with these new resins a treatment of one minute only at 120° C. is sufficient; the textile material being treated is thus less liable to impoverishment.

The following is an example of the manner in which viscose rayon fabric is given a good feel and made uncrushable; First a solution is prepared with the following ingredients:

30% Formaldehyde	900 lb.
Urea	300 lb.
30% Ammonium Sulpho-	
cyanide Solution	150 lb.
Water	900 lb.

The fabric is impregnated with this liquor, squeezed free from excess, and then dried. Afterwards the fabric is led over rollers heated to about 130° C. and the impregnated substances then react to form an elastic insoluble resin which makes the viscose fibers practically uncrushable.

It is possible to use an ammonium sulphide instead of the more expensive sulphocyanide and also to color the fabric during impregnation with the resin components. Thus viscose rayon fabric is impregnated with the following liquor:

	30% Formaldehyde	900 lb.
	Urea	300 lb.
	30% Ammonium Sulphide	150 lb.
	Sulphonated Cetyl Alcohol	
	(Wetting and Dispersing	
	Agent)	60 lb.
	Ammonium Sulphocyanide	50 lb.
	Diamine Sky Blue FF	20 lb.
u	ad then dried at 150° C. for	10 minutes

Crease Resisting Fabric U. S. Patent 1,980,676

Fifteen gallons of casein solution containing 1 lb. of dry casein and 2 oz. of trisodium phosphate are mixed with 5 gal. of 30% latex solution containing 2% zinc oxide on the dry rubber and 2% piperidine penta-methylene dithiocarbamate. The latter material acts as an accelerator An ordinary sizing for the rubber. mangle can be used, the excess size being removed and the fabric is then dried. Subsequently, the fabric is washed in boiling soap solution to remove that part of the size which held the latex in suspension, presumably the casein compo-In order to prevent the crossed yarns from adhering to one another, work the fabric during the drying operation which is the method employed in the acid organdie process for the same purpose.

Delustering Finish for Rayon

3. Sulphonated Castor

- Fuller's Earth
 Titanium Dioxide
 40 lb.
- Oil (30%) 150 lb. 4. Stearic Tallow Softener 15 lb.

Mix 1 and 2 and wet out with 3. Then add 4 and grind well.

Degumming and Decolorizing for Straw British Patent 424,189

Soda Ash	80	lb.
Rosin	80	lb.
Casein	250-300	lb.

Water to give consistency of soft soap while being heated.

Renovating Surfaces of Textiles British Patent 419,856

The shine produced on textile fabrics by wear can be removed if the fabrics are first dry-cleaned, the surface fibers raised by teazelling, and then a mixture of 1 part sodium salicylate, 2 parts borax, 1 part cresol saponatis, and 3 parts ammonia in 320 parts water applied; finally the goods are brushed thoroughly.

Mercerizing Wetting Out Agent U. S. Patent 2,008,458

Cresol, Technical	90	lb.
Aniline	10	lb.

Mercerizing German Patent 606,025

As wetting agents for use in mercerizing lyes, use is made of acid esters of phosphoric acid in association with phenols and (or) highly sulphonated oils. A typical wetting agent comprises dibutyl phosphate 1, crude cresol 9 and a highly sulphonated oil 2 parts by weight.

Low Luster Artificial Silk U. S. Patent 1,967,206

Casein		10	lb.
Water		200	lb.
Turpentine		10	lb.
Petrolatum			

10% of weight of cellulose
The above is emulsified and added to
the spinning solution (viscose).

Partially Saponifying "Celanese"

To dye directly and uniformly with certain dyes, it is often necessary to par-

tially saponify "Celanese" by padding with the following and drying.

Soda Ash 30 lb. Glycerin 2 gal.

After drying, steam for 4 minutes in a rapid ager. Rinse well and dye with any direct dyestuff.

Restoring Luster to "Celanese" Pad with 28% acetic acid, tenter and dry under tension. Rinse well and dry.

Rejuvenating Cloth U. S. Patent 2,006,192

A composition suitable for treating worn shiny wool or silk fabrics is formed of alcohol 16 oz., 24% ammonia solution 3 oz., glacial acetic acid 4 oz., oil of lavender 1.5 g. and chloroform 2 oz.

Cotton Softener

a. Tallow 4 g. Caustic Potash (50° Bé.) 1.2 g.

When a is saponified, add b with stirring and stir until solidification begins.

Pre-Shrinking Treatment of Cotton Fabrics

U. S. Patent 1,959,406

Cotton fabric is shrunk by immersion for 1-10 hours in an aqueous liquor at 65-100° containing 1-4 oz. of ammonium alum and 0.25-3 oz. of sodium bisulphate per 10-50 oz. of water, followed by hydro-extraction (without intermediate washing) and drying.

Tarnish-Proof Cloth U. S. Patent 1,933,302

The cloth after dyeing is dipped in a solution of a cadmium salt (0.5 lb. or gal.) e.g., cadmium acetate which absorbs hydrogen sulphide when used as a wrapping for copper and silver articles and thus protects them from atmospheric tarnishing.

"Cravenetting" Textiles

The process of waterproofing or cravenetting proper is not a simple one. Soaking the fabric in a strong solution of acetate of alumina for several hours, extracting and allowing to dry slowly, is about as effective as any simple process. The acetate of alumina may be prepared by dissolving 1 lb. of alum in 1 gal. of hot water. In another vessel containing

½ gal. of water dissolve 1¼ lb. of sugar of lead (lead acetate). Mix the two solutions and allow the precipitate to settle. The clear liquid only is used in preparing the bath, using about 1 qt. of the solution to 1 gal. of water.

Proofing Against Moth and Fungi British Patent 413,445

Animal fibers such as wool, felt, fur, skins, feathers, silk and hair, are proofed against moth and fungi by treatment with a solution of chromium fluoride so that a definite quantity of chromium compound equivalent to 0.65% of chromium fluoride is retained by the material. After steeping or padding with the aqueous solution, excess is removed and the chromium compounds fixed on the fiber by drying at a temperature above 150° F. In British Patent 413,529, the process in the above specification is modified by adding antimony fluoride to the chromium fluoride bath.

Mould and Fungi Proofing of Textiles British Patent 413,648

About 5% barium borate is claimed as an impregnant.

Silk Wool for Knitting

Silk-wool, suited for knitting, may be produced as follows: The woolen yarn is first treated for 15 to 30 minutes in a cold bath of 100 l. in which 34 l. of hydrochloric acid (at 32° Tw. == 1.160 sp. gr.) has been dissolved. The yarn is now to be well drained or else hydroextracted. A second cold bath is prepared by using the clear liquor from a solution of 1½ kg. of bleaching powder in 100 l. of water. The yarn is treated in this cold bath for perhaps 15 to 30 minutes. Afterwards the yarn is drained and then soured with hydrochloric acid for 30 or 45 minutes. Next, the work is to be rinsed and then turned for 15 to 30 minutes in a warm bath at a temperature of 75° C. (167° F.). This bath is to contain 600 g. Marseilles soap per 100 l. of water. The work is now removed and hydro-extracted. Afterwards, it is given a second souring with hydrochloric acid. Finally, it is well washed.

Felt Hat Stiffener

Carnauba Wax Emulsion (Bright Drying) Shellac (Ammonia Water Solution)

90 lb.

10 lb.

Stiffening Material for Shoes French Patent 777,404

The material is made by impregnating cloth, paper or felt with a colloidal substance, a part of which is in the precipitated state and consequently easy to dissolve while the rest is not precipitated and therefore less easy to dissolve. Thus, flannel is impregnated with a colloidal solution containing cellulose nitrate 150 kg., alcohol 580, acetone 60, carbon tetrachloride 120 \(^1\). and then dipped in water for 15 minutes. A part only of the nitrate is precipitated and the material is air dried.

Rubber Latex as a Textile Finishing Agent

The use of rubber as rubber latex or in a dispersed form has found many applications of late in the textile industry. It is natural to assume that a substance possessing the characteristics of rubber, i.e., water repellency and its flexibility, and especially the fact that it may be applied to a textile in a liquid state like many other finishing compounds, should find development in the finishing of textiles.

The application of rubber latex in connection with textiles has been grouped as follows: For the production of artificial leather and non-skid rug underlaps; as a backing and sizing for pile fabrics, or binding and strengthening agent for fabrics that otherwise would be too sleazy for rough usage; for double texture fabrics. Hauser has discussed the use of latex in combination with canvas for friction belts, as well as its use as a binding agent for applying flocked wool or cotton to a fabric base.

The utilization of rubber latex in the carpet industry has assumed a rôle of importance as carpetings impregnated with it form their own selvedges without unravelling, thus obviating the necessity of a binding. Carpetings of this type may be joined together by use of a latex adhesive without any evidence of a surface seam. If the proper latex is used for the backing of the carpet, the latex is waterproofed to such an extent that it may be scrubbed on a floor without the moisture coming through. Rubber latex has been an important factor in developing a new type of construction in carpets and pile fabrics. In this process, a hair batt is laid on a latex-coated base and the fabric subjected to a vulcanizing process. In this particular development the use of looms for the production of the carpets has been done away with entirely.

It has been stated that it is obvious that the textile mill is not equipped to develop the various latex compounds required. A textile plant possessing the facilities of the average sizing and finishing equipment and laboratory will probably be in a position to develop rubber latex as a finishing agent.

Rubber Latex

The presence of rubber latex as a processing agent has been made possible because of developments in prolonging its stability. Crude rubber latex, when stabilized with ammonia immediately following tapping, will withstand reversion or coagulation for the interval of ship-ping time until it reaches its destination, where it is subjected to further stabilization with ammonia. The rubber latexes are white to grayish in color, and are found occasionally with a yellowish cast. Latex, when freshly collected from the tree, may contain as high as 50% rubber, but following stabilization the rubber content will drop usually to 40% and under. In a number of cases, before selling, it is concentrated by various methods, or is compounded for a particular need.

The concentrating of latex is carried out by various processes, which may be subdivided as follows: (1) by creaming promoted by centrifugal force much in the same manner as a cream separator; (2) by filtration through unglazed porcelain while the latex is kept in movement; (3) by evaporation after the latex is stabilized by a non-volatile stabilizer like soap or sodium alginate.

Water dispersions of rubber differ from latex in that the latter at no time in its processing has been reverted to the solid state, but has been kept liquid since its tapping from the tree. The water dispersion, on the other hand, is a stable dispersion of coagulated, smoked rubber, plus various compounding ingredients, effected by mechanical means. have been marketed by a number of the leading rubber companies already compounded, and they exhibit properties similar to rubber latex towards other chem-They are usually less expensive than latex and greater efficiency may be obtained by their use because of the greater rubber concentration of the majority of dispersions when compared to the ordinary 40% latexes. Water dispersions of rubber usually yield softer films, but one of their drawbacks lies in that many of these are not as lightly colored as rubber latex and consequently will not yield the latter's clear films.

Rubber Latex with Starch

Rubber latex may be incorporated with a starch sizing to add flexibility and water resistance when padded to a fabric. Crude latex in admixture with a starch sizing will not waterproof a fabric but it will enhance its water repellency. However, if a compounded rubber latex is used, waterproofedness will be produced.

Rubber latex in mixture with starch is used extensively today as an adhesive. The mixture is not an easy one to produce. This is due to the action of a starch paste, which, although it is itself a protective colloid, tends to coagulate latex when it is added in a hot state. The latex should be first protected with a protective colloid such as glue, casein or gum tragacanth. Bone glue has been found to be an excellent protective agent as well as one exhibiting properties akin to a starch.

One part of a better grade of bone glue is heated, while stirring in 8 parts of water, to 140° F. until all lumps have been dispersed, and a smooth thin paste results. The glue should not be heated to over 140° F. since a decomposition of the protein may result.

If the cooked glue is tested for acidity it will be found to be somewhat on the acid side. Any substance exhibiting an acid reaction should not be added to rubber latex as acidity will tend to coagulate it. Consequently the glue is made alkaline with 0.5% solution of caustic soda, and cooled to about 110° F. (Although precautions against the addition of caustic soda to latex have been advised, no deleterious effects from the addition of small amounts of it have as yet been noted.) The latex—four parts of latex to one part of glue by volume—is then further stabilized with a small amount of ammonia, and then poured slowly while stirring into the glue. Thus we now have the protected latex mixture.

The starch (maize cooked 1 lb. to 1 gal.—tapioca starch 8 oz. to 1 gal.) paste is cooled to 140° F. and an equal volume of water is added. This should be made alkaline with a small amount of ammonia; the protected latex mixture is added to it slowly and stirred until a uniform mixture results. If this size mixture is padded on to a cotton fabric, a firm, flexible finish will result. Thus in a like manner it may be thinned to yield the desired firmness.

In adding a protected latex solution to

a cooked starch, care should be taken that the size should not be too hot-not over 140° F., since there is a liability of coagulation of the latex. Once a latex reverts or coagulates, there is little hope for its redispersion, since this may be carried out only with special equipment as that used for making water dispersions of rubber. However, there are certain indications of partial coagulation before a latex will completely revert. If, upon the addition of the protected latex to the starch, a sudden stiffening of the latter is noted, we have an indication that coagulation is setting in. No further addition of latex should be made, but the starch should be further thinned with ammoniated water until it thins out evenly, and then the remainder of the latex is added slowly.

A size-latex mixture as prepared above will produce a water-repellent finish on a fabric but will not waterproof it. To produce a waterproof finish, a "curable" or vulcanizable latex must be used.

Compounding Rubber Latex

In order to compound crude latex for vulcanization, there are certain essential chemicals which should be present in the mixture at all times. These are sulphur, zinc oxide, and an accelerator. Sulphur chloride may be substituted for sulphur. Any other chemicals added are for the purpose of lending some desired property to the resultant rubber film.

Any substance added to latex must be water-soluble and completely miscible with it, in order to produce effective results. Sulphur and zinc oxide in their dry state are not soluble in water and therefore cannot be incorporated into latex as such. Sulphur chloride is miscible with latex, but because of its cost and its irritating action on the skin should be disregarded. Thus the zinc oxide and sulphur must be placed in a water-soluble state before their addition to latex. This is done by placing them in a colloidal state, and they are marketed as colloidal sulphur and zinc oxides and capable of being thinned to a great extent with water before they fall out of solution. On a dry basis, the concentration of dry sulphur in the colloidal material is about 45% by weight, while the zinc oxide runs about 54% by dry weight.

The purpose of the sulphur in the mixture is to produce greater flexibility and toughness in the rubber film. To hasten this effect, zine oxide is added. It may be termed a very slow accelerator in the vulcanizing or "curing" action of the sulphur on the rubber. However, to

hasten the reaction between the rubber and sulphur to a greater degree, a more rapid outside accelerator is invariably added as well. Water soluble accelerators are present on the market which will cause the rubber to vulcanize at a temperature of 140° F., and it has been noted that latexes compounded with these accelerators vulcanize at oven temperature. A simple starting recipe for a vulcanizable rubber mixture is:

 $\begin{array}{cccc} \text{Latex (50\%)} & 1 & \text{gal.} \\ \text{Colloidal Sulphur (45\%)} & 11 \text{4} & \text{oz.} \\ \text{Colloidal Zinc Oxide (54\%)} & 2 & \text{oz.} \\ \text{Accelerator} & 1 \text{2} & \text{oz.} \\ \end{array}$

In preparing this mixture, the colloidal sulphur and zine oxide are first thinned separately with a portion of the latex before their addition to the major portion. The accelerator is first pasted with a little sulphonated castor oil, and then thoroughly dissolved in a small amount of water at 150° F. The solution is then strained through a cheese cloth into the partially compounded latex. The latter is then stirred thoroughly to produce a uniform mixture.

If an accelerator which must be emulsified before adding to latex is used, it should be emulsified with triethanolamine

and oleic acid as follows:

Accelerator	100 lb.
Oleic	5 lb.
Triethanolamine	2 lb.
Water	80 lb.

The accelerator and oleic acid are thoroughly mixed and added slowly while stirring to the triethanolamine diluted with the water. The amount of this emulsion added to the latex should be based on the actual weight of the accelerator present in a specific volume. For liquid accelerators, the dispersing of these in water with ammoniacal casein is recom-An agitator must be used in order to obtain a stable dispersion. In order to prevent rubber films from oxidizing too rapidly, compounds called anti-oxidants are often incorporated into the latex batch. For the majority of water-soluble anti-oxidants used with latex, the amount used is about double the weight of accelerator in the formula. If this vulcanizable mixture is protected with glue in the same manner as the crude latex and then added to a size batch which is applied and dried into a fabric, a complete waterproof should re-

Care should be taken in drying fabrics impregnated with a starch-crude latex sizing on a can dryer. A crude latex film

when subjected to heat has a tendency to become soft and sticky, thus tending to adhere to the dry cans. If the percentage of latex in the size batch is such that this occurs, the sticking may be overcome by powdering the cans with a small amount of talcum. With a tenter dryer, little difficulty should be encountered in this direction.

In coating fabrics with latex for adhesive purposes or for producing protective films, it is desirable that greater amounts of latex should be carried to the material. This is accomplished by use of a thickening agent on the same principle as the use of a thickener in printing fabrics. A more concentrated latex may be used alone since it is naturally creamy and thick. A natural 40% latex, however, must be thickened. Thickening agents include starch, water-soluble resins and colloidal clays. Where a coating is desired which overlooks the brittleness produced by the starch, then the latter should be used. Colloidal clays should be used when the natural flexible rubber films are sought. Much of the firmness as produced with a starch may be overcome by the addition of a softener such as sulphonated castor oil. If an excess of the sulphonated castor oil is used, tackiness in the crude rubber film re-

Of the clays, a good grade of colloidal bentonite makes an excellent thickening A concentration of 1 lb. to a gallon of water in admixture with 1 gal. of crude latex yields a viscosity which produces continuous films having good To produce the clay paste, the dry bentonite should be first pasted with a small amount of sulphonated castor oil thinned with a portion of the subsequent water to be used. The remaining water is then stirred in and the mixture allowed to soak overnight for the lumpy clay to expand. On the following day the paste is thoroughly mixed and then strained through cheese cloth before its addition to be used. The remaining water is then stirred in and the clay will tend to dust when it is found present in the rubber film.

If it is desired that the film should be colored, an organic dye in solution may be added, but the greatest fastness is obtained by use of water-soluble dispersed colloidal pigments which are present on the market.

Films produced from crude latex mixtures, as pointed out previously, will tend to grow tacky with heat. If this condition is undesirable, a compounded latex must be used.

gal.

Wetting Agents with Latex

Recently, a number of wetting agents have been marketed especially for use with latex. These are of use when a thorough impregnation of a heavily woven cotton fabric is necessary. A wetting agent showing an acid reaction when in solution should be avoided. The best method of accomplishing a thorough impregnation of a heavy cotton fabric is first to boil it out thoroughly in soda and in a wetting agent, and after a thorough wash and nipping it should be run through a pad in open width containing the latex and wetting agent.

If the material is but wetted in water and the wetting agent added to the latex bath, then the high speed of the pad should be diminished. Instead, the cloth in open width is run very slowly through the latex in order to insure a thorough soaking, and then through the nip.

Precautions in Handling Latex

- (1) There should be a word of advice to the workman handling latex, and this is that he should abstain as far as possible from placing his hands in the raw latex. The reason for this is that in many cases there is an acidic reaction from the perspiration on the hands which tends to cause reversions. Cases of latex coagulation have been reported due to this cause.
- (2) Rinds and latex films that are noted on the surface of a latex bath should be picked off, since these hasten congulation. If possible, when these occur, the bath should be strained through a cheese cloth to remove the films.
- (3) Latex should not be subjected to abnormal conditions of temperature. Latex when frozen will coagulate when reliquefied, and consequently should never be stored in a spot where a low temperature of 32° F. may occur. Latex should not be heated as this will cause the stabilizing ammonia to volatilize, this condition tending to hasten coagulation.
- (4) Latex mixtures should not be made in copper vessels, since if small amounts of copper are present in a rubber film the metal will tend to hasten the oxidation of the film.
- (5) Latex should never be added to size baths containing calcium, barium, or aluminum salts, as these exert a coagulation action on latex.

Rubber latex has found a place for itself in the finishing of certain textiles. It can be handled properly with the finishing equipment of the average mill. The prime requisite is that the finisher

familiarize himself with this somewhat new finishing agent.

Fireproofing Solutions

The following is the formula of a solution used in theatrical work for rendering materials non-inflammable:

Tungstate of Sodium $17\frac{1}{2}$ oz. Water $1\frac{1}{2}$ pt.

Dissolve in the cold and add:

Sodium Phosphate 2½ oz. Water 1 pt.

or a sufficiency of water to make the solution sp. g. 1.140.

Dip the material in the solution, wring out with the hands, dry, and iron if nec-

The following are formulæ of solutions advised by the L.C.C. for rendering curtains, Christmas decorations, etc., non-inflammable:

Formula No. 1

Ammonium Phosphate	1	lb.
Ammonium Chloride	2	lb.
Water	$1\frac{1}{2}$	gal.
No. 2		Ŭ
Borax	10	oz.
Boric Acid	8	oz.

Water

Both solutions can be used for coarse fabrics, but No. 2 is better for more delicate articles. The fabrics should be dried without rinsing, and it is advisable to experiment with a small portion of the cloth before treating the whole, as the texture and colors of some materials are affected detrimentally.

Fireproofing for Canvas

Ammonium Sulphate	8	oz.
Ammonium Carbonate	2.5	oz.
Boric Acid	3	oz.
Borax	2	OZ.
Starch	2	oz.
Dextrin	0.4	oz.
Water	100	oz.

Steep ½ hour at 86° F.; 2 dips necessary for best results.

Fireproofing Brake Lining U. S. Patent 2,001,194

Brake lining is impregnated with a composition such as may be formed from an aniline dye 10 to 20 g., ammonium sulphate 60 lb., ammonium phosphate 10 lb., boric acid crystals 12 lb., gum acacia 2 lb., cresley ore 2 lb., barium hydroxide 4 lb., aqueous ammonia 1 qt., ammoniumaluminum sulphate 2 lb., copper-sodium

alginate 1.5 lb., benzaldehyde 1 oz., sodium bicarbonate 2 lb. and water 100 gal.

Flameproofing and Fireproofing Textiles
Sodium Borophosphate Resin
(Abopon) 8 lb.

Water 5-6 gal.

Dip the textile into the above solution warmed to 110 to 170° F.; wring out and pass between warm rollers. This process gives a uniform coating which does not powder out like the usual fireproofing salts.

Waterproofing Canvas Formula No. 1

A treatment that is sometimes given to awnings to waterproof them and still leave them flexible so they can be rolled up and down, is as follows: First apply a coat of glue size, made by dissolving 1 lb. of high-grade glue in 3 qt. of water. To 1 gal. of this size add 1 oz. of alum, previously dissolved in hot water. Apply the size while still quite warm, using a wide flat wall brush. When the size is dry apply two coats of a paint made by mixing white lead-in-oil, with necessary tinting colors added, thinned to rather stout brushing consist-ency with a liquid composed of 2 parts of boiled linseed oil and 1 part of turpentine. Be sure to use boiled linseed oil, as raw oil would have a greater tendency to rot the canvas, more especially if glue size has not been used under the Two coats, or not more than 3 coats, should be sufficient. Be sure to allow ample time between coats for thorough drying. If the use of paint is objectionable, shave paraffin into gasoline, in the proportion of 2 oz. of paraffin to 1 gal. of gasoline, stirring until the wax is dissolved. The wax must be in very thin shavings to dissolve quickly in cold gasoline. soon as the wax is dissolved, brush a coat of the solution on the bare canvas, using a wide flat wall brush. The next day another coat may be applied. If you brush the material on carefully you should be able to build up a reasonably smooth, waterproof surface in this way. Be very careful when using this preparation that no one strikes a match near you, and that there is no sort of flame in the room where you are using the solution, or you may have an explosion.
One of these processes embodies the use of paint and the other a wax as the

waterproofing agent, and either will leave the canvas reasonably flexible and waterproof.

No. 2 Canvas Waterproofing

1 0		
Gilsonite	10	lb.
Asphaltum	2	lb.
Degras, Neutral	4	lb.
Beeswax Crude	1	lb.
Lead Oleate	3	lb.
Kerosene	31	lb.
Gasoline	41	lb.

Waterproofing Cotton Cloth

Pad the cloth with aluminum acetate solution (2° Tw.) and dry. Then immerse in sodium stearate "solution" (5%) at 120° F. Rinse well and dry.

Tarpaulin or Tent Waterproofing Formula No. 1

British Patent 414,242

Paraffin Wax	3-5	lb.
Naphtha	200	lb.

Warm together on steam bath and mix until clear. Then mix in:

Aluminum Powder 5–20 lb.

No. 2

Australian Patent 17,598

Rubber Latex	1-2 lb.
Linseed Oil	½ lb.
Casein	2 lb.
Water	16 gal.

Water-Repellent Fabric U. S. Patent 1,967,267

Fabric is impregnated with a solution of 1 pt. of wax (or animal and vegetable fats, greases, or oils) and 1 pt. of water shedding substance (e.g., cellulose acetate or nitrate, etc.) in an organic volatile solvent (e.g., ethyl acetate) and then dried, whereby it retains its original softness but becomes water repellent.

Textile Backing (Waterproof)

Latex (50% Concentration	1) 1	gal.
Casein	12	oz.
Water	1	qt.
Zinc Oxide	$1\frac{1}{2}$	OZ.
Sulphur	5/8	oz.
Accelerator No. 552	1/2	oz.
Agerite White Powder		
(Anti-Oxidant)	5/8	oz.

Waterproofing Wool Goods

The simplest method of waterproofing wool goods is the application of metallic

salts and tannic acid, sold either as powder or crystallized, with or without previous or subsequent soap, or fatty acid baths.

Formula No. 1

For 100 l. of impregnation bath there is dissolved about 100 g. of acetate of lead, 200 g. of alum, and 100 g. of tannin in boiling hot water. The goods repassed at about 40° C., centrifuged, and dried at from 40 to 50° C. The effect of the impregnation process is considerably increased by the above-mentioned soap and fatty acid baths.

No. 2

Three hundred grams of the best sulphonated oil, and 100 g. of olive oil soap are stirred in 10 l. of boiling water. They are added to a bath of 90 l. water at a temperature of 50° C. and the goods are passed at 40° C. To simplify the procedure these two baths may be combined in one.

No. 3

One hundred grams acetate of lead, 200 g. alum, 100 g. tannin, 20 g. linseed oil, 500 g. Monopol oil, and 100 g. of pyridine are well stirred into about 20 l. of boiling water and brought to a boil again. Then the whole is increased to 100 l. by adding water of at least 60° C. The goods are passed at 40° C. and dried rapidly. Wool fat that can easily be emulsified is also well suited for the wet impregnating of wool. When it is used, the emulsifying is done separately.

No. 4

Ten kilograms of wool fat, 1 kg. ammonia, 5 kg. sulphonated oil, and 500 g. pyridine are brought to the boil in about 50 l. of water, the whole being well stirred. This suffices for an impregnating bath of about 800 l. Into this bath, before adding the emulsifying agent, there are stirred 500 g. of pyridine, and the temperature is brought to 50° C. The goods are dipped at from 30 to 40° C., centrifuged, and dried thoroughly. make the impregnation more effective, there may be added to these baths tannin substances or metallic salts. effect is always superior when they are used in separate baths.

Waterproofing Wool, Silk, Rayon and Cotton

Examples for impregnating fabrics and wearing apparel of wool, silk, rayon and cotton are as follows: In 100 l. of petrol or other volatile hydrocarbon solvent, are dissolved by stirring well, 1

kg. of linseed oil varnish and 2 kg. of ceresin, the latter first being melted. The goods are thoroughly dipped, centrifuged, and dried in the open air. Subsequent steaming gives further assurance of even and thorough impregnation throughout the fabric. Fabrics can be steamed on a wet pressing roller. With very light colored and with white goods, the best wool fat is used instead of the linseed oil, and white paraffin instead of ceresin. Wool fat is recommended especially for wool goods when a soft feel is to be preserved, since after the admixture of varnish, the goods grow harder with time. The varnish impregnation is particularly suitable for coarser goods for which very thorough waterproofing is desired, especially for tentings, army blankets, water pails, and for colored umbrella fabrics of all kinds of fibers.

Porous Cloth, Waterproofing

For this purpose a solution of acetate of alumina or acetate sulphate of alumina, which is prepared as follows, is chiefly used.

Sulphate of Alumina 665 lb. dissolved in

Water 600 lb. Sugur of Lead 945 lb. dissolved in

Water 900 lb.

Dissolve each by itself hot, precipitate cold, draw the clear solution off and make to Twaddell 15°. In this manner, a standard alumina sulphate-acetate is obtained of which the greater part is deposited on the fiber in drying.

As woolen and half wool goods still contain some soap from the milling process, a soap passage is as a rule not necessary before impregnating with alumina; otherwise the goods are passed through a weak soap solution (3:1000), squeezed and dried without rinsing.

The goods are impregnated on a hank washing or open width washing machine provided with pressure rollers, by passing the dry goods for 1 hour through the diluted acetate-sulphate of alumina of 3% Tw. (undried goods at 7½-15° Tw.). The goods are then slightly centrifuged without rinsing or squeezed and then dried.

For wool and half wool goods a single impregnation will suffice in most cases: if a higher grade of waterproof finish is desired the treatment is repeated, inserting a soap passage if necessary.

In place of acetate-sulphate of alumina, formate of alumina may be used

with advantage. The latter possesses the advantage over the former that the danger of the subsequent tendering of the cotton warp in half wool goods, due to the formation of sulphuric acid in the fiber, is eliminated. Formate of alumina is used in the same manner as acetate-sulphate of alumina.

Waterproofing and Fireproofing Fabrics, Paper, etc.

Austrian Patent 136,953

The material is coated or impregnated with an alcohol solution containing a resin, fat or like substance and a non-hydrolyzing salt of a metal of the 2nd periodic group which forms a colorless or transparent compound with the alcohol. A typical solution comprises resin 2, castor oil 0.5, crystalline zinc chloride 3, crystalline magnesium chloride 5, and 96% alcohol 12 parts. The solution may be applied to crepe paper.

Waterproofing and Flameproofing U. S. Patent 2,003,148

A method of compounding a composition of matter for flame and waterproofing aqueous cellulose media and their derivatives comprises heating 640 parts of water to 120° F., adding 48 parts of ammonium sulphate and stirring until completely dissolved, adding 16 parts of ammonium carbonate incrementally under constant stirring until effervescence ceases, adding 20 parts of boric acid previously dissolved in 128 parts of boiling water, adding 16 parts of borax and thoroughly mixing, adding 16 parts of starch previously cooked to about 1° Bé. and thoroughly mixing in the same under constant stirring; dissolving 6 parts of suitable soap in 128 parts of water and bringing it to the boiling point, thereafter adding the same to the previously compounded materials, bringing about emulsification of the whole and then lowering the temperature to about 110° F. and digesting for about 2 hours thereby forming a first composition; bringing 640 parts of water to the boiling point and dissolving therein 80 parts of ammonium chloride, 48 parts of boric acid and 16 parts of borax in the order named, and each after the preceding has been completely dissolved, stirring the same thoroughly after all three have been added and dissolved, separately dissolving 32 parts of soft gelatin in 256 parts of water and heating to about 200° F. under constant stirring and

thereafter optionally adding thereto 131/2 parts of glycerin, stirring thoroughly and then adding the same to the ammonium chloride-boric acid-borax solution under constant stirring for about 30 minutes and then digesting for about 1 hour at about 140° F.; dissolving 3 parts of suitable soap in 128 parts of water, heating to boiling and adding 8 parts of dextrin, stirring such constantly to insure uniformity and then adding such to the ammonium chloride-boric acid-borax-gelatin solution, thereby forming a second composition; bringing 128 parts of water to the boiling point, dissolving therein 15 parts of soap bark and filtering, thereby forming a third composition; dissolving 32 parts of alum in 256 parts of water as a fourth composition; digesting each of the four compositions for about 4 hours while stirring from time to time; combining the first, second and fourth compositions in a common vessel and then adding the third composition under vigorous stirring.

Colloidal Textile Oil

rolling ro. r	
Castor Oil	20 gal.
Coconut Fatty Acids	100 gal.
Caustic Soda Solution	•
(30° Bé.)	15 gal.
Water	30 gal.

Manipulation: Mix in the order given at 40° C.

Castor Oil 15 gal. Coconut Fatty Acids 75 gal. Water 22½ gal. Caustic Soda Solution (30° Bé.) 11½ gal. Paraffin Oil (28° Bé.) 82 gal. Manipulation: Mix at 40° C.

Colloidal Olive Oil Commercial Olive Oil Caustic Potash Solution (32° Bé.) Water 13 lb. 150 lb.

Manipulation: Stir the caustic potash solution into the olive oil at room temperature and allow to stand overnight. In the morning add the water (which is previously brought to a boil). The mixture is well stirred during addition of the water, which is added slowly.

	Acetate Rayon	Oil		
	Sulphonated Castor Oil			
,	(65%)		50	gal.
	Commercial Olive Oil		45	gal.

Acetic Acid	(28%)	20 gal.
Paraffin Oil	(28° Bé.)	5 gal.
Water		100 gal.

Manipulation: Mix the three oils and the water at 40° C. Then cool to 30° C. and stir acetic acid into mixture slowly.

Hosiery Oil

Sulphonated Castor Oil		
$(\bar{6}5\%)$	1000	lb.
Caustic Soda Solution		
(27° Bé.)	300	lb.
Water	650	lb.
Manipulation: Mix caustic	soda	solu-
tion with oil at 40° C., then	add	water
slowly, maintaining temperat	ure a	t 35-
40° C.		

Kier Penetrant Oil

Xylol	10 gal.
Sulphonated Castor Oil (62% T.F.M.) Water	20 gal. 20 gal.

Manipulation: Sulphonate the castor oil to 62% T.F.M., settle and draw off. Mix in xylol first and then water, with agitation, at 35-40° C.

Silk Oil

Sulphonated Castor Oil	
(58%)	50 gal.
Paraffin Oil (28° Bé.)	10 gal.
Caustic Soda Solution	Ü
(27° Bé.)	12 gal.
Water	23 gal.
Steam Distilled Pine Oil	12 gal.

Manipulation of Silk Oil: Mix ingredients in order named at 35-40° C., being careful to add caustic soda solution and pine oil very slowly, with constant stirring and allowing mixture to cool to room temperature as the pine oil is being added.

Soluble Oil Formula No. 1

Paraffin Oil (28° Bé.)	33	gal.
Sulphonated Castor Oil (75%)	22	gal.
Sulphonated Red Oil (75%)	33	gal.
Manipulation: Mix at 40° C.		

3.7	_
No.	2

Steam Distilled Pine O	il 50 gal.
Sulphonated Castor Oil (75%)	50 gal.
Caustic Soda (27° Bé.)	10 gal.
Water	40 091

Manipulation: Heat the pine oil to 38° C. in the lead lined tank, add the

sulphonated castor oil, then add the caustic soda gradually with agitation, maintaining the temperature noted above with constant agitation. When nearly classolution is obtained add the water slowly, continuing agitation, then allow to cool rapidly.

Soluble Textile Oil

Manipulation: Mix the paraffin oil and red oil, heat to 40° C., add the previously mixed water and caustic solution, then add the xylol slowly and the alcohol last and rapidly cooling them as quickly as possible after mixture is uniform.

Wool "Soluble" Oil U. S. Patent 1,965,935

An oil such as a mineral oil 64, is used in admixture with "Carbitol" 2, corn oil soap 14, rosin 10, water 6 and diethylene glycol 4%.

Wool Treating Oil Formula No. 1

Neutral Light Mineral Oil Double Pressed Red Oil		gal.
No. 1 Lard Oil		gal. gal.
Manipulation: Mix at 45-50°	C.	0

Equipment required: Wooden or lead lined mixing tank.

No. 2

Paraffin Oil (28° Bé.)	90	gal.
Double Pressed Red Oil No. 1 Lard Oil		gal.
Manipulation: Mix at 45-50°		gal.

Textile Sizing Oil

167 dig dig ou		
Sulphonated Castor Oil		
(62% T.F.M.)	800	
Water	550	lb.
Caustic Soda Solution		
(27° Bé.)	350	lb.
Silicate of Soda Solution		
(37° Bé.)	1300	lb.

Manipulation: Heat the sulphonated oil to 35-40° C. and slowly add the other ingredients in order given above, maintaining temperature above 35° C. until mixing is completed.

Oiling for Viscose Yarn

Ammonium Oleate Oleic Acid	$100 \\ 25 - 30$	
Alcohol Apply at 40-60° C.	15	ğ.

A 1% solution of above works well at 40° C.; treating time 25 to 30 minutes.

Rayon Yarn Lubricant U. S. Patent 1,979,188

Mineral Oil Sulphonate Potassium Oleate ''Carbitol''	9.7 9 6 5	lb. lb. lb.
Aniline	0.3	lb.

Synthetic Neat's Foot Oil

Extra Lard Oil	30 gal.
No. 1 Lard Oil	30 gal.
Light Mineral Oil	30 gal.
Manipulation: Mix at 40°	C.

Rayon Identification (Revised Method)

The following systematic scheme, when carried out in the given sequence, serves for the rapid identification of rayons. This method can be depended upon by an experienced analyst, particularly when used in conjunction with filament count and microscopical characteristics. For the inexperienced man we recommend the detailed method of Rayon Analysis, and in comparison of the unknown rayon with standard samples of known make. The standards should be as inclusive of the rayon field as possible and should be kept up to date.

Rapid Method

Test 1-Identification of Animal Fibers

Millon's Test

Animal fibers—real silk, wool and hair—are quickly and positively identified by means of Millon's Reagent (see Identification of Rayon—Detailed Method).

Test 1A—Identification of Animal Fibers, Cellulose Fibers and Cellulose Acetate

Flame Test

Twist five or six strands of the un known sample into a long, compact mass. Push the end of sample gently toward a match flame. (Do not allow sample to actually touch the flame.)

Animal fibers tend to fuse and burn slowly when brought near to a flame. If the flame of the burning fibers is extinguished, the odor of the white fumes which subsequently arise from the smoldering end will have a ''burned hair'' odor. The burned ends of the fibers will have a dark, hard, brittle knob of material. Heavily mineral-weighted silks will leave a distinct ash which more or less retains the shape of the original material.

Vegetable fibers and most rayons do not fuse in the burning. They burn rapidly, and the fumes coming off after the extinguishing of the flame smell like burning cotton. Acetate rayons, in burning, smell like cotton and melt like animal fibers. The fused knob remaining after the flame is extinguished is hard but not brittle. If heated to a sufficient degree (in an evaporating dish or other suitable container) acetate fibers will melt without burning.

The burning test, while helpful, is not as instructive as the Millon's Reagent Test, inasmuch as it does not show the relative quantities and locations of the animal and vegetable fibers in mixed varns or fabrics.

Test 2—Identification of Cellulose Acetate Rayon

Solvent Test

Cellulose Acetate Rayon is soluble in acetone; also in boiling 40% acetic acid. (See Identification of Rayon—Detailed Method.)

Test 3—Identification of Nitrocellulose Process Rayon

Diphenylamine Test

Nitrocellulose Rayon is turned blue by treatment with a solution consisting of 1% by weight of diphenylamine dissolved in concentrated sulphuric acetic acid mixture. (See Identification of Rayon—Detailed Method.)

Test 4—Identification of Viscose and Cuprammonium Rayons

Wright's Stain Test

Wright's Stain Test solution colors air-dried Cuprammonium Rayon violet and air-dried Viscose Process Rayon blue. (See Identification of Rayon—Detailed Method.)

Detailed Method

Chemical Identification of Rayon

(1) Identification of Animal Fibers in Mixed Fabrics

Millon Test

As small quantities of animal fibers present in unknown samples may cause

confusion in some of the following tests, an unknown sample should first be tested for the presence or absence of animal fibers. These are easily and quickly identified by means of the Millon Test, details of which follow:

Preparation of Millon's Reagent

Millon's Reagent is prepared by dissolving a given weight of metallic mercury in its own weight of pure concentrated nitric acid at room temperature in a non-corrodible container (porcelain, glass, agate, etc.). When completely dissolved, the solution is diluted and mixed with an equal volume of cold water. The solution should be clear.

If a yellow turbidity develops in the above noted solution, stir in a small quantity of nitric acid until the solution clears

up.

Each new batch, or one which has stood open to the air for a long time, should be tested for proper activity by matching to the skin or by use of white animal fibers. When stored in air-tight glass stoppered bottles, the solution keeps for months.

Use of Millon's Reagent

Moisten the unknown swatch with Millon's Reagent. Warm to blood heat (do not boil) for a few seconds, or allow to stand for a few minutes at room temperature.

Animal fibers turn red, thus showing both their presence and position or distribution throughout the pattern.

Nearly all dyed animal fibers show an observable change toward red in this test without previous stripping of dye.

Swatches wet with water or with alcohol appear to react normally if flooded with reagent (to dissolve first precipitate).

Caustic Test

As minute quantities of cellulose and rayon fibers present in unknown samples largely composed of animal fibers may not be detected by the Millon Test, we recommend a subsequent caustic test for fabrics that appear by the Millon Test to be composed largely or entirely of animal fibers.

Although strength of solution, time and temperature may be varied over wide limits we recommend a 10% solution of caustic soda at 180° F. for 10

minutes.

Animal fibers dissolve completely. Cellulose and rayon remain in fiber form. (Note-Cellulose Acetate is partially saponified; regenerative cellulose fibers soften and dissolve to a limited extent. Cellulose Acetate fibers may be removed previously to caustic boil by use of acetone.)

(2) Identification of Cellulose Acetate Rayon

(a) Acetone Test

Place yarn or fabric in U.S.P. ace-

Cellulose Acetate is very readily dissolved.

So-called "iron-proofed" Cellulose Acetate partially dissolves and leaves a leathery residue.

All other Rayons are unaffected by this treatment.

(b) Acetic Acid Test

Place yarn or fabric in a boiling solution of 40% acetic acid (C.P. acid is not necessary).

Cellulose Acetate is very readily dis-

solved.

So-called "iron-proofed" Cellulose Acetate partially dissolves and leaves a leathery residue.

All other Rayons are unaffected by this treatment.

"''Iron-Proofed" Cellulose Acetate "Iron-proofed" Cellulose Acetate is Cellulose Acetate that has been treated with an alkaline medium in such a way that the outside of each individual filament is partially samparified

ponified. "Iron-proofed" Acetate yarn may be pressed or ironed at a higher temperature than un-treated acetate, because the layer of saponi-fied or partially saponified acetate insulates the unaffected core of the yarn.

Treatments similar to iron-proofing, but more drastic, produce partially saponified yarn

or fabric that can be dyed with direct dyes. Partial saponification other than for ironproofing is occasionally practiced. Such yarns produce a very considerable residue when reated by the acetone or acetic test for Cellulose Acetate.

(3) Identification of Nitrocellulose Rayon

Apply one drop of diphenylamine solution* to the dry unknown sample.

Nitrocellulose Rayon immediately turns a deep blue color after which it slowly dissolves to form a blue solution.

Other rayons are not colored blue. All nitrated fibers—for example, Viscose Process Rayon nitrated for the production of special effects-show a blue reaction with diphenylamine solution. Many dyestuffs show a blue coloration when exposed to diphenylamine solution.

Nitrocellulose samples that have been stripped in a strong reducing bath will sometimes fail to give the blue coloration

* Diphenylamine solutions is prepared as follows: Mix 66 g. concentrated sulphuric acid with 33 g. of glacial acetic acid, then add 1 g. diphenylamine.

described above, however, their crosssections remain unaltered in shape.

The only positive test for Nitrocellulose Rayons is a microscopic examina-

(4) Identification of Viscose and Cuprammonium Rayon

(a) Wright Stain Test

Prepare a saturated solution of Wright Stain (dry powder) in denatured alcohol (95%). Immerse air-dried unknown sample into boiling Wright Stain solution and boil for a few seconds. Rinse the sample thoroughly in cold water.

Viscose Process Rayon is stained blue

by this treatment.

Cuprammonium rayon is stained violet. Schreiber-Hamm (Sulphide) Test

This test is suitable only for raw rayon of standard manufacture. Certain experimental yarns and processed yarns cannot be positively identified by this

A 5-g. sample of the unknown rayon (Viscose or Cuprammonium) is placed in a flask together with 100 cc. of water and 3 cc. concentrated sulphuric acid. The mouth of the flask is covered with a piece of lead acetate paper and allowed to stand on a moderately boiling steam bath for 4 hours.

If the sample is Viscose Process Rayon, the lead acetate paper will be stained

brown or black.

If the sample is Cuprammonium Rayon, no discoloration should be observed.

(5) Identification of Undesulphurized Viscose Process Rayon

The difficulty of visually distinguishing between some delustered rayons and undesulphurized Viscose Rayon has sometimes led to confusion and improper rayon identification.

Undesulphurized Viscose Process Rayon can be readily identified by means of so-

dium plumbite solution.

Preparation of Sodium Plumbite Test Solution:

- (1) Dissolve 40 g. lead nitrate in 200 cc. of warm water.
- (2) Dissolve 70 g. of caustic soda in 300 cc. of water.
- (3) Add the caustic soda solution to the lead nitrate solution.
 - (4) Filter.
 - (5) Dilute to 2 1.

Method of Testing

A small quantity of the solution prepared as above is brought to the boil.

The unknown rayon sample is inserted into the boiling test solution for a period of 1/2 minute.

Undesulphurized viscose process yarn

turns black.

Incompletely desulphurized viscose process yarns are turned black, dark brown, or medium brown, depending on the degree of desulphurization.

Desulphurized viscose process yarn is

stained a brownish yellow color.

When possible, check tests on known samples should be run simultaneously with the test.

Microscopic Identification of Rayon

As rayons are most easily, quickly and positively identified by means of a microscopic examination, this method should be used whenever possible.

A microscopic examination of rayon is very simple and can be successfully carried out by men previously unfamiliar with the use of the microscope after a

few hours' practice.

For the benefit of those unfamiliar with the microscope and its use, we are pleased to describe the cheapest type of microscope that is, in our opinion, suitable for the microscopic examination of rayon. The analyst will need:

Microscope Stand and Lenses.

The instrument should be capable of magnifying to 400 diameters.

The above combination includes achromatic objectives, 16 mm. and 4 mm., eye piece $5\times$ and $10\times$; and Abbe condenser N.A. 1.20.

Microscope Lamp.

3. Microscope Slides andGlasses.

 Mounting Medium (Methylene Iodide, or Monobromnaphthalene).

A piece of thin glass rod.

6. A small scalpel or sharp knife.

Treatment of Viscose Products Austrian Patent 138,007

Rayon and other products made from viscose are bleached and desulphurized by treatment first with an alkaline solution of hydrogen peroxide at a low temperature and then with an alkaline solution not containing hydrogen peroxide at a raised temperature. Thus, rayon may be treated at atmospheric temperature with a solution containing hydrogen peroxide 0.5 and sodium pyrophosphate 1%, freed from excess of liquid, left to stand for 3 hours at 35° C., and then treated at 95° with a solution containing sodium pyrophosphate 1 and Marseilles soap 1%. Alternatively, the material may be

treated with a single alkaline hydrogen peroxide solution first at a low temperature and later at a raised temperature.

Preservation of Ropes

Make a solution of sulphate of copper (blue vitriol) in water, using 1 lb. of the crystals in 4 gal. of water and soak the ropes in this solution for 4 days, then dry them. The ropes will become impregnated with the copper sulphate, which will keep them from being attacked by parasites and prevent rot. The copper salt may be fixed in the ropes by the application of a soap solution, made by slicing 1 lb. of yellow laundry soap in thin slices and dissolving it in boiling water. Use 1 lb. of soap to a gallon of water. While the soap solution is still lukewarm put the ropes in it and let them soak overnight. Next morning take the ropes out and let them dry. The copper soap thus formed is more effective than tar, which is used on ropes employed by sailors, but tar is likely to stain painted surfaces, so painters should stick to the soap treatment. Ropes must be kept in a warm, dry place, never in a basement, because dampness would injure them in

Sash Cord Impregnants Formula No. 1

Paraffin Wax	130-		
132°F. M.P.		8	0 z.
Rosin		4	oz.
Rosin Oil		1	oz.
Carnauba Wax		1	oz.
	No. 2		
Lactic Casein		10	oz.
Borax		2	oz.
Pigment		60	oz.
Soap Solution		4	oz.

0.5 oz.

0.8 oz.

Caustic Soda

Ammonium Sulphate

Water remainder
The soap solution can be sodium resinate or the potassium salt formed by boiling potassium carbonate (1 part) with carnauba wax (15 parts). The ammonium sulphate is added after all the other ingredients are in solution. The pigment could be china clay or tale colored to shade with a brown lake. The composition given would require further adjustment with water to give the right consistency in the coating tank.

Numida Dyeing of Feathers

Dissolve gum arabic in cold water to about the thickness of varnish.

Make up a solution containing:

Gum Arabic Water 1 glass Cold Water 2 glasses Glycerin 1 glass

Strain thoroughly to remove all particles of dirt, etc.

Take the dry feathers and work in this solution until thoroughly saturated, wring through the ordinary wash wringer, and squeeze out as much of the solution as possible, after which rub through the hands thoroughly for about 5 minutes in order to evenly distribute the remaining portion of the liquid that is in the feathers, after which string the feathers and beat them out on a wooden board for several minutes until the fine stems separate, after which hang up and dry overnight.

Feathers thus treated will retain this effect under all ordinary conditions.

Fabric Paint

Basic Dye	2	lb.
Ethylene Glycol	60	lb.
Zinc Chloride	6	lb.
Tannic Acid	6	lb.
Glacial Acetic Acid	6	lb.
Tragacanth Solution (1%)	90	lb.

Synthetic Resin for Impregnating Textiles

British Patent 422,957

Polyvinyl Chloride (60-		
65% Chlorine)	5-10	lb.
Methylene Chloride	6	lb.
Benzene	3	lb.
Butyl Acetate	1	1h

Weighting Cotton Yarn

Cotton yarn may be weighted to a considerable extent, when dyed with the direct colors, by adding magnesium sulphate (Epsom salt) to the dye bath, together with a small quantity of dextrin. Owing to danger of imperfections in the color, such as unevenness and cloudiness, it is perhaps better to use a separate bath after the dyeing for the purpose of weighting. This will be especially true if it is desired to weight to any considerable extent. The following process is a typical example of weighting cotton yarn which has been dyed with direct colors. For 100 lb. of cotton yarn use a bath containing about 160 gal. of water; add 100 lb. of magnesium sulphate, 15 lb. of dextrin, and 2 lb. of glycerol. Have the temperature of the bath at about 120° F. The cotton yarn is entered into this bath and turned for 20 minutes, or until the fiber is thoroughly saturated with the solution. It is then removed, hydro-extracted and dried. Such a treatment as this will give a weighting of about 10 to 12% to the cotton yarn. The bath is by no means exhausted, and may be freshened up by the addition of a small amount of magnesium sulphate and dextrin till it is brought back to the same hydrometer test as at first, and succeeding lots of cotton may be treated as above. The glycerol is added for the purpose of preventing the weighting material from giving the fiber a stiff handle.

Rayon Spinning Solution

To a solution of 25 parts acetonesoluble cellulose acetate and 75 parts of

95% acetone plus 5% water is added 2.5 parts of a mixture containing mineral oil (100 viscosity at 100° F. Saybolt) 85, saponifiable oil (olive oil) 10, tetrahydronaphthalene 2.5 and soap (sodium oleate) 2.5%. The yarn spun from the solution is bright and fairly transparent and has superior knitting properties.

Wet Strength of Wet Fibers, as a Percentage of Their Dry Strength

Cotton	110-120%
Wool	80- 90%
Silk (True)	75- 85%
Acctate Silk	65- 70%
Cuprammonium Silk	50- 60%
Viscose Silk	45- 55%
Nitro Silk	30- 40%

MISCELLANEOUS

Boiler Compounds

Formula	No.	1
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Sodium Alginate (Crude)	20 lb.
Extract, Quebracho	12 lb.
Soda Ash	10 lb.
Trisodium Phosphate	10 lb.
Caustic Soda	1 lb.
Water	300 lb.

Manipulation: Dissolve the salts in the water and add the alginate and quebracho extract at room temperature.

No. 2

Anhydrous Disodium	
Phosphate	47 lb.
Soda Ash	44 lb.
Corn Starch	9 lb.

It should be noted that this formula includes both inorganic and organic constituents. The starch is added to bring about a state of colloidal suspension of the insoluble matter precipitated in the boiler so that a sludge is formed in preference to a scale.

Another composition which deserves consideration is the U. S. Navy Standard Compound, which is:

No. 3

Anhadaana Cadima Canhanata	70	112
Anhydrous Sodium Carbonate	10	ID.
Trisodium Phosphate	10	lb.
Dextrin or Starch	1	lb.
Cutch sufficient to yield	2	lb.
tannic acid		
Water to make up to 3	100	lb.

Coal Dust Briquettes German Patent 616,376

Finely divided coal sludge brought to water content of 12 to 20% is mixed with 2 to 3% molasses and then compressed in molds and dried.

Fuel Briquettes for Motors

One hundred kilograms of sugar or molasses are mixed with 5 kg. of alum or a similar substance for inversion of the sugar and dissolved in 400 to 600 kg. of water, after which finely ground bituminous coal is added until a homogeneous mixture is obtained. The mixture is

poured over 50 to 100 kg. of a finely disintegrated mass of sugar beets. Thirty to 50 parts by weight of the mass thus obtained are mixed with 70 to 50 parts of finely ground charcoal, and the mixture is pressed to briquettes under a pressure of 100 to 300 kg. per sq. cm. The briquettes are dried by heating in a separate drying chamber by means of combustion gases from a steam boiler furnace. The drying requires only about 15 to 30 minutes, during which the briquettes take on a cokelike appearance. Owing to the high temperature in the drying chamber, about 350° to 500° C., and the high water content of the briquettes, steam is formed during the drying which seems to have a hardening effect upon the briquettes. Under this high drying temperature the sugar content of the briquettes is caramelized. A suitable composition of the dry matter of the briquette mass is stated as 80 parts by weight of charcoal, 20 parts of bituminous coal, and 2 to 6 parts of sacchariferous binding substances.

Fuel Briquettes U. S. Patent 1,977,332

Slowly burning briquettes suitable for use in orchard heaters are formed by mixing charcoal 50, sand 25 and a sugarsyrup binder about 25% so that all the particles of charcoal and sand are coated by the syrup, molding without applying pressure, evaporating moisture from the briquette in the mold and then heating to about 370° C. for about 2 hours to form an anhydrous porous mass, and cooling under air-tight conditions.

Briquettes

French Patent 766,979

Semicokes and fine coals are mixed with 6-12% of pitch, molded and heated to about 600° C. and then carbonized at 700-900° C.

Battery Paste

In the manufacture of lead-acid storage battery plates it frequently happens that the paste in the plates checks when

dried. The addition of a small amount of silicate of soda to the paste will reduce this tendency. The amount should be not over 1 oz. of the strong solution of silicate of soda (water glass) to 100 lb. of the oxide. This should be dissolved in about 1 pt. of water and added to the oxide before the acid is added.

Low-Voltage Storage Battery Paste U. S. Patent 1,944,065

The paste for a lead accumulator contains (a) 0.9 to 1.5 weight per cent of nickel sulphate, or (b) 0.1 to 0.5 weight per cent of cobalt sulphate as active material.

Cold Storage Fluid U. S. Patent 1,943,268

Fluids for cold storage comprise water (in each case) and butyl alcohol 10%, or ethyl ether of glycol acetate 20, or diethylene glycol butyl ether 5%.

Low Freezing Heat Transfer Medium U. S. Patent 1,972,847

A stable heat transfer medium comprises a mixture of 60 parts of diphenyl oxide, 12 parts of naphthalene, 28 diphenyl.

Antifreeze Composition Formula No. 1

A mixture of 65% isopropyl and 35% methyl alcohol is recommended for addition to radiator water. It does not attack the metal parts and changes the boiling point of water only slightly.

No. 2

U. S. Patent 1,997,735

A cooling medium having a freezing point below -45° F. and a boiling point above 217° F. consists of a solution formed by adding 2 lb. of calcium chloride and 7 oz. of aluminum chloride to glycerin, 1 pt., and water as 1 gal.

Prevention of Ice Formation on Airplanes U. S. Patent 2,017,593

A mixture of liquids of different effects on rubber (such as pine oil 4, diethyl phthalate 4 and castor oil 1 part) is used in such relative proportions as not substantially to swell or otherwise deteriorate a rubber surface to which the composition is applied.

Anti-Knock Fuel Formula No. 1

U. S. Patent 2,021,088

0.5 to 5% of ethylene diamine or 0.5 to 1% of a hydrate of the same is used with gasoline.

No. 2

U. S. Patent 1,973,320

A mixture for introduction into the cylinders of internal combustion engines to prevent knock or pinking and the deposition of carbon comprises 85 g. of uranium chloride and 15 g. of vanadium chloride dissolved in acetone.

No. 3

U. S. Patent 1,980,097

Chloral hydrate in small quantities may be utilized to assist the solution of the metallic chlorides. For example, 1 to 10 mg. of platinum chloride may be dissolved in 1000 cc. of butyl oxalate and 250 to 2500 mg. of vanadium chloride in the same amount of butyl oxalate. The solutions are then combined and sufficient butyl phthalate is added until it constitutes about 25% of the mixture.

Stabilization of Anti-Knock Compounds British Patent 414,581

Decomposition of lead tetra-ethyl present in the fuels is prevented by the addition of a small amount, e.g., 0.01—0.05% of sodium fluoride, potassium fluoride or ammonium fluoride.

Detergent for Automobile Radiators Formula No. 1

U. S. Patent 1,967,393

A mixture is used comprising ammonium hydroxide or cyclohexanol 1, a sulphonic acid derivative of an alkylated aromatic hydrocarbon such as sodium 1-isoprioylnaphthalene-2-sulphonate about 0.4 and an alkali metal carbonate such as sodium carbonate about 4 parts.

No. 2

U. S. Patent 1,967,394

This relates to a detergent mixture comprising an organic solvent immiscible with water such as ammonium hydroxide or cyclohexanol about 1, a sulphonic acid derivative of an alkylated aromatic hydrocarbon about 0.4 and sodium phosphate about 4 parts.

Carbon Electrodes for Batteries British Patent 429,840

A mixture of finely-ground bone charcoal 34, wood charcoal 8, graphite 6, pinewood flour 8, ammonium sulphate 14, and sulphur 6 parts, with a binder made by stirring a mixture of wheat flour 6, sugar 18, water 7.5, and oil 15 parts at 80° C. for 15 minutes to burst the starch granules and dissolve the sugar, is extruded or pressed into the desired electrode shape, dried, and fired in cast iron boxes or in saggers packed in graphite or retort-gas carbon, the temperature being raised slowly to 1000° in 16 hours and maintained there for 4 hours. After cooling, 1/4 of the block is immersed in a 2-5% solution of paraffin wax in petrol and the other 34 is then immersed for 3-4 minutes in 10% aqueous ammonium chloride. The waxed top is then drilled, a copper terminal screwed in, and the joint again waxed. Finally the whole electrode is impregnated with a 10-12.5% solution of silicic acid in trichloroethylene, carbon tetrachloride or other volatile solvent and dried.

Brake Fluid Composition U. S. Patent 1,928,956

A hydraulic fluid comprises, in solution, glycol acetate, e.g., 50% by volume, with smaller proportions of water 37-45 and sulphonated castor or linseed oil soap, 5-13.

Moisture-Resistant Bristles U. S. Patent 1,953,980

The bristles are first impregnated with an aqueous heavy-metal salt (e.g., 1-3% aqueous aluminum acetate) and then with a water soluble soap of a fatty acid (e.g., 4% aqueous castile soap). They may also be dipped into a solution of a wax in xylene.

Catalyst Canadian Patent 350,894

To a dry mixture of kieselguhr 150, gum tragacanth 10 and potassium sulphate 20 lb. is added with agitation a sodium vanadate solution prepared by treating 16 lb. of vanadium pentoxide with 10 gal. of water containing 11.3 lb. of sodium hydroxide. The mixture is diluted with 20 gal. of water and after thorough mixing sulphuric acid is added to neutralize or nearly neutralize the mixture. The mixture is evaporated to a yonsistency suitable to permit granula-

tion or pelleting and the granules or pellets are heated for 1 hour at 600° C. The product is a catalyst for the oxidation of sulphur dioxide.

Catalyst for Ammonia Oxidation U. S. Patent 2,017,683

Metallic cobalt, containing impurities 70, is heated to effect fusion with calcium carbonate 3.5–5 and calcium fluoride 1.7–3.5 parts, the slag formed is separated from the metal and the latter is converted into cobalt oxide.

Activation of Kaolin for Catalytic Purposes

Kaolin is ignited at 750-800° C. for 2 to 3 hours and treated in the cold with 33% nitric acid for 24 hours and the solution is then heated at 60-80° C. for 3 to 4 hours. Aluminum hydroxide is then precipitated and allowed to stand for 1 day at room temperature before filtration. It is dried at 100 to 120° C. and activated at 360-385° C. The catalyst is suitable for the dehydration of alcohol.

Regeneration of Spent Nickel Catalysts

The method consists essentially in treating the spent catalyst successively with a small quantity of 20° Bé. sodium hydroxide, sulphuric acid and water. Before saponifying the spent catalytic mass, it is heated with indirect steam with vigorous stirring till a homogeneous mass is obtained. The sodium hydroxide solution (60-80 l. for 500 kg. of catalyst) is then added, followed by sufficient water to make the mass fluid; saponification is effected by heating with stirring for 11/2 to 2 hours. After transferring the soap to a lead-lined tank, it is decomposed with concentrated sulphuric acid, diluted with water and allowed to stand, and the supernatant fat is removed. The nickel is then boiled with sulphuric acid as usual. The recovery of nickel is 92-94%, as compared with 64-70% by the ordinary method.

Fuel Catalyst French Patent 765,824

A mixture used for activating the combustion of solid fuels contains, e.g., manganese dioxide 32.1, organic material (wood charcoal) 2.5, sodium chloride 27.7 and sodium chlorate 37.7%.

Cable Insulation U. S. Patent 1,946,322

The mixture comprises a hydrocarbon oil (e.g., cylinder oil) 95-50, and rosin free from oxidized components, especially abietic acid, 5-50%.

U. S. Mint Test Solutions for Counterfeit Coins

Gold

Concentrated Nitric
Acid
Hydrochloric Acid
Distilled Water

Concentrated Nitric
6½ drachms
5 drachms

Silver

Silver Nitrate 24 gr.
Nitric Acid 30 drops
Distilled Water 1 oz.

A drop of the above solutions will have no effect on genuine coins; but will stain others, i.e., spot them.

Capsules British Patent 412,975

Capsules or coverings, for bottles, jars, metal tubes and rods, of the kind made from a composition containing cellulose ester and a substance which may be removed by a suitable solvent after formation of the capsule, etc., to cause the capsule, etc., to shrink on drying and fit tightly onto the article to which it is applied, are formed by compression, extrusion or injection from a composition produced by working or mixing together the cellulose ester, a water soluble softener and optionally, a plasticizer to produce a solid but plastic composition. Small amounts of a volatile solvent may be In an added to facilitate mixing. example, a mixture containing cellulose acetate 3, monochlorohydrin 2, monoglycerol benzoate 1 and water 2 parts is mixed at 80-100° C. until completely gelatinized and most of the water has evapo-The material is then formed to the desired shape and rendered contractile by soaking in water to dissolve out the monochlorohydrin. The contractile capsule is then applied to the article on which it is to be used and, as the water dries out, the capsule shrinks into position. Filling materials, dyes or pigments may be added.

Motor Carbon Remover U. S. Patent 2,004,628

A carbon removing composition is composed of kerosene, creosote, castor oil and

amyl acetate, combined in substantially the following proportions: kerosene, $49\frac{1}{2}\%$; creosote, 25%; castor oil, 25% and amyl acetate, $\frac{1}{2}\%$.

Activating Adsorbent Clay U.S. Patent 1,976,127

The method of activating adsorbent earths comprises mixing an earth with concentrated sulphuric acid in an amount equal to from 5% to 35% of the weight of the earth, heating the mixture to a temperature of 150 and 300° C. to obtain reaction of sulphuric acid with constituents of the earth and to also partially dry the earth and the products of such reaction by the combined effect of heating and the dehydrating action of the sulphuric acid, then bringing the resultant mixture into contact with water to dissolve soluble salts therefrom, separating the solution from the undissolved earth, and then drying the earth.

Processing Coal Canadian Patent 324,976

Coal containing iron sulphide is thorroughly washed to remove dust and impurities and while wet is sprayed with a compound containing calcium chloride 92, potassium dichromate 3, manganese dioxide 3 and tannic acid 2 parts by weight. The burning properties and ash characteristics are improved and the deleterious effect of flue gases and tube-slagging is minimized.

Oil Treatment of Coal U. S. Patent 2,005,512

The process of treating solid lump fuel to render the same dustless, consists of heating oil having a gravity of 19° to 30° Bé. at 60° F. and a Saybolt viscosity of 100 to 1200 at 100° F. to a spraying temperature of 100° to 250° F., and spraying the heated oil in finely atomized state on the fuel in quantities sufficient to deposit on the fuel a thin enveloping film of oil.

Fuel Oil Activator Japanese Patent 101,701

Thirty grams of the above is added to 5 gal. fuel oil to increase heating efficiency.

Dustproofing of Coke

A 1 to 1 emulsion of a thick petroleum oil and water is made at 94° C., and then diluted with 7 parts of water at 38° C. Two gallons are sprayed per ton of coke on the loading chutes.

Decolorizing Charcoal from Corncobs

Soak corncobs in 3% zinc chloride and 7% sulphuric acid for 24 hours. Distill destructively at 600° C. for 50 minutes and treat with superheated steam at 400° C.

Deodorizing Petroleum

Petroleum products may be conveniently deodorized by agitating thoroughly with quicklime, 3 oz. to the gal. and filtering.

Gasoline Gum Inhibitor U. S. Patent 1,970,339

Nicotine pyrogallate or amylgallate is added in proportion of about $\frac{1}{100}\%$.

Coloring Leaded Gasoline Canadian Patent 352,875

 $\alpha(\mbox{2-Methoxyphenylazo})\mbox{-2-naphthol}$ is used at rate of 2 to 12 oz. per 10,000 gal.

Liquid Dielectric Composition U. S. Patent 1,999,004

Chlorinated biphenyl having a chlorine content of 60% is used in a proportion of 45% together with trichlorobenzene 25 and tetrachlornaphthalene about 30%.

Condenser Dielectric

A 50% solution of Bakelite in castor oil has a high dielectric constant, 5.6, as compared with 2.3 for transformer oil. A condenser having paper impregnated with the Bakelite mixture has a power factor of 1% against 0.5% with transformer oil, this being the only disadvantage.

Dielectric Materials

French Patent 765,876

Dispersions of metal soaps in insulating oils are used, e.g., 6-10 g. of aluminum stearate in 94-90% of oil.

"Coreth" Type A	rtificial	Diesel	Fuel
Alcohol		36	kg.
Coal-Tar Oil			kg.
Gas Oil		20	kg.
Wood Oil		10	kg.
Water			kg.
Degras, Saponified	i	2	kg.

Liquid Electric Insulation British Patent 413,596

A mixture comprising mineral hydrocarbon oil (50-70 parts) and halogenated diphenyl, e.g., the polychlorinated derivative (50-30 parts).

Electrical Insulator British Patent 429,730

Rutile	32 lb.
Talc	58 lb.
Blue Clay	6.5 lb.
Calcium Carbonate	3.5 lb.
Mix thoroughly and mold.	

Electric Insulating Compositions German Patent 616,056

A binder for use in making insulating compounds or materials comprises a resin, a vegetable drying oil, shellac and (as a flux) an aromatic compound boiling above 200° C. A specified binder comprises copal 12.5, wood oil 1.5, α -nitronaphthalene 1 and shellac 10 parts. Mixtures of the binder with subdivided mica or like material may be molded under heat and pressure, or a solution of the binder in an organic solvent may be applied to mica sheets and the latter then united by heat and pressure.

Electrical Insulating Fused Magnesia British Patent 413,905

The electrical resistivity of fused magnesium oxide is permanently increased by heating slowly to 1149° F., maintaining it at this temperature for about 6 hours and finally cooling to room temperature in about 30-40 hours.

Vitreous (Electrical Insulating) Material

U. S. Patent 1,984,178

Silicon dioxide is fused with beryllium oxide 0.14-1.5 and aluminum oxide 0.2-2.0%.

Waterproofing Electrical Wires Formula No. 1

Crepe Rubber Mineral Spirits	30 lb.
Mill together until uniform,	
Glue Solution (20%) followed by	25 lb.

20 lb.

Water

	No. 2				1
U. S. Pa	tent :	1,963	,895		4
Mineral Oil				70 cc.	60
Neat's Foot Oil	ĺ			25 cc.	cle
Ethyl Acetate				1 cc.	Co
					cha
Electrical :		_	Tap	е	Bla
	ula I				
Make up caoute			ions.		
a. Caoutchouc,	Crude	, in		00 1	
Smoked Pres b. Benzene or 1	sea s	neets	3	20 kg. 80 kg.	
	No. 2	ше		ov kg.	
Resin (_				!
Formu		b	c	d	;
Rosin Rosin Oil	$\frac{40}{36}$	30 30	20 28	20 g. 30 g.	1 :
Rosin Tar	30	10	40	30 g. — g.	
Petroleum Tar			10	— g.	80
Stearin Tar			10	— g.	
Coal Tar				20 g.	1
Wood Tar,					
Anhydrous		7.0	_	10 g.	
Mineral Oil Linseed Oil	$\frac{-}{24}$	$\frac{10}{20}$	$\frac{8}{24}$	10 g.	1
				10 g.	o arc
The formulae at the two others. It pale resins, as a, a	anu For w	υ ar hita	e su ribb	perior i	v con
pale resins, as a.	are po	ssibl	e.	ons, om	8%
	No. 3	,00101	٠.		an
		D:			diz
Fillers For White Ribbon		r ign		7.	rap
Lithopone	18		<i>a</i> 80	b 60 g.	
Zinc White			20	60 g. 20 g.	
Barium Sulphat	е		_	20 g.	
For Black Ribbon	ıs		c	ď	
Barium Sulphat			40	20 g.	anı
Vegetable Black	:		45	g.	of
Lamp Black			15	15 g.	str
Frankfort Black Chalk Powder	ζ			45 g.	pre
	No. 4		_	20 g.	box
Definit			ra		00.
White Cover Ribb		ıı.xıu.			
Rubber Solution	(No	1)		38 g.	
Resin Compositi	on (I	No. 2	a)	22 g.	
Fillers (No. 3a,	3b)		/	40 g.	
Black Ribbon:			a	b	
Rubber Solution	(No	. 1)	44	42 g.	
Resins (No. 2b, Resins (No. 2d)	2c)		26	_ g.]]
Resins (No. 2d)	9.71		20	30 g.	ber
Fillers (No. 3c,			30	28 g.	
	No. 5				
Coating to	Be A	ppli	ea b	У	
Hot Ir	чħьев	ша і 10	иT]
White Coating	(000	a s		E7	sil
a. Linseed Oil Crude Rubbe	(0U° '	U.) Smr	11	57 g.	litt
a. Crude Rubbe		\sim ma	**		exp

Piecer

	Resin						g.	
c.	Filler	S	$(N_0$. :	3a, 3b)	28	g.	
Pr	epare	α	in	a	kneading	machin	e :	a

Prepare a in a kneading machine at 60° C., then heat up to 180° C.; when clear solution is formed, add melted b. Cool to 100° C., and add c, stir, and discharge above 70° C.

0			
Black Coating			
Linseed Oil		60	g.
Crude Rubber		6	g.
Rosin Tar		12	g.
Fillers (No. $3c$,	3d)	22	ğ.
	or		
Mineral Oil		52	g.

 Crude Rubber
 6 g.

 Petroleum Tar
 12 g.

 Wood Tar, Anhydrous
 10 g.

 Fillers (No. 3c, 3d)
 20 g.

 Thèse masses should be kept at 60

These masses should be kept at 60-80° C. in the impregnation vat.

Fusible Cut-Outs British Patent 423,076

A fuse wire incorporated in a current consuming device, e.g., an incandescent or arc-discharge lamp, rectifier or valve, is composed of a brass containing 0.25 to 8% aluminum, e.g., copper 67, zinc 32, and aluminum 1%. This is non-oxidizable, has a higher resistance and melts rapidly.

Electrolytic Condensers British Patent 421,628

An electrolytic condenser is wound in annular form to permit free circulation of air around it. Aluminum electrode strips are separated by strips of cloth impregnated with an electrolyte, of the composition glycol 400 cc., borax 25.6 oz., boric acid 17.0 oz. and water 25.6 oz.

Electrolytic Condenser Medium U. S. Patent 1,973,554

Monoethanolamine	1 lb.
Ethylene Glycol	5 lb.
Boric Acid	5 lb

Heat together until dissolved and add bentonite or starch to consistency desired.

Fingerprint "Raising" from Cloth

Dip in, or paint with a 10% solution of silver nitrate to which has been added a little acetic acid. Dry in dark room, then expose to ultra-violet light until of maximum intensity, and photograph.

Latent Fingerprinting

A piece of paper or other material on which one is searching for fingerprints is saturated in a sensitizing solution prepared by dissolving 2 g. of silver nitrate in 1 l. of distilled water. This is stored in a dark place. After having soaked for 2 hours in the silver nitrate bath, the paper is thoroughly washed in distilled water, first by soaking for 30 minutes and then two rinsings. There is left in the paper only the silver chloride which has been formed from the chlorides left by the perspiration and the silver nitrate. The paper is hung up and allowed to dry thoroughly. It is then developed, either with a developer of the M. G. type or with others, such as formaldehyde and sodium carbonate. Following the development the paper is again washed in water, then in a bath of hypo, washed, and dried, and is ready for observation.

If kept in a humid atmosphere the migration of the chlorides may be so intensified that in time a gray cloud is formed where the print was originally. In some cases the print goes through the paper. Prints made from the skin of a corpse are very poor and diffuse, although

chloride is deposited.

Fire Extinguisher U. S. Patent 2,010,729

A fire extinguishing composition comprises 48 parts by weight of sodium bicarbonate, 12 parts by weight of boric acid, 4½ parts by weight of potassium bitartrate, and about 1½ parts by weight of borax.

Fireproof Film Containers British Patent 419,249

The walls are made of a mixture of sawdust 25, calcined magnesite 25, magnesium chloride (as a 25% aqueous solution) 30, potassium alum 10 and a mixture consisting of asbestos flour 4, asbestos fiber 3 and acetic acid 3, 10 parts.

Fluorescent Screens French Patent 770,728

The screen contains zinc or cadmium borate, e.g., zinc silicate 10-12, calcium tungstate 45-50 and zinc or cadmium borate 40-45%.

Electrotyping Matrix British Patent 430,660

A sheet of aluminum 0.007 in. thick, is cleaned with etching fluid or caustic

soda and then coated with molten beeswax, preferably a mixture of gum damar 2 and beeswax 16 parts, heated to 160° F. The wax face is then coated with graphite to render the surface conductive.

Masking Taste of Chlorinated Water Add 2 or 3 tablespoonfuls of wine to each liter of water.

Fish Baits

The common "baits" comprise two general categories: (1) artificial baits,

and (2) natural baits.

Artificial baits may be classified as flies, spoons, spinners, phantoms, and a multitude of other contrivances, some of which may be used alone, and some in combination with natural baits. Flies are largely made of feathers, worsted, silk, tinsel, etc., and are fashioned to suggest an insect. Most flies, however, resemble only remotely any known insect. Other baits may be made of metal, wood, rubber, etc. The list is too extensive to enumerate here, and more may be learned from a reliable fishing tackle dealer than from reading pages of descriptions. With reference to natural baits, with which the following lists are concerned. a local angler can usually impart to the novice more practical knowledge in a short time than could be learned from a whole volume of discussion and descriptive matter.

Judging by the stomach contents of fishes, there are but few groups of animals, from worms to mammals, that do not afford food for one or another game fish. That some of these are occasionally swallowed by a fish, however, does not necessarily signify that they would make good bait. Furthermore, some of the best baits can never be the natural food of the fish. The groups of animals which comprise forms most commonly employed as bait, from the lowest form up, are: worms, mollusks, insects, crustaceans, fishes, birds, and mammals.

It must be borne in mind that baits used in one part of the country may be of little or no avail in another part, even for the same species of fish; and that in the same locality the proper baits often vary with the time of year. Furthermore, a killing bait of one day may prove ineffective on the next. Success in fishing, therefore, depends largely upon the experience, judgment, skill, and patience of the fisherman.

Vernacular names of the various animals used for bait differ greatly in dif-

ferent parts of the country; for instance, the stone fly of one section is the mill fly of another, and the hellgramite of one locality is the dobson of another, and so on. Therefore, any list of baits can be of only partial assistance. The following lists aim to give the most common baits under the names by which they are most widely known.

Natural Baits are used in several different ways, such as in still fishing, bait easting, skittering (modified form of easting), or trolling.

Live Bait.—It has always proved practically impossible to keep a large amount of live bait in restricted limits; furthermore, no fish will live indefinitely without food. The kind of food necessarily depends upon the kind of fish, but most shiners and other minnows are more or less carnivorous and finely ground meat of some kind would probably answer for this class. The most appropriate food, however, would be small crustaceans and aquatic insects such as are usually present in sluggish streams and small ponds. These may be collected by means of a gauze dip net. It is possible to stock a small pond or pool, or even a rain barrel, with small crustaceans and maintain a supply of that kind of food. Some species of bait minnows are much hardier than others, but in all cases, when kept in confinement much depends upon the maintenance of cleanliness and a sufficient supply of oxygen. The needed oxygen is best supplied by a continuous flow of well aerated water, but where this is impossible it may be furnished in a fine spray of compressed air introduced near the bottom of the tank. Cold water will dissolve more oxygen than warm water, therefore, the temperature should be kept low if possible. Overcrowding should be carefully avoided and all injured or sick fish should be removed as soon as detected. If feeding should be attempted great care should be taken to remove all food uneaten, as otherwise it will decay and pollute the water.

Conditions will vary according to the species of minnow, the size and character of the tanks or pools, the temperature of the water, and the number of fish per unit of space, and it is difficult, therefore, to furnish specific information without a knowledge of these factors.

Keeping and Rearing Earthworms for Bait.—Earthworms multiply by producing eggs which are laid in capsules in the ground. The young become fully grown in four or five months. One method of culture is to sink into the soil

in some shady spot a box of suitable size, usually not less than 18 inches deep and of any desirable width. The top of the box should be made hinged, or removable, and placed from two to three inches below the surface of the surrounding soil. This box should be nearly filled with rich, dark loam that should be kept quite moist but not wet, for too much water will kill the earthworms quickly. The worms may then be collected and placed in this box, and may or may not be covered with a layer of green sod.

By far the easiest and most convenient way to collect earthworms is by the use of a flashlight or lantern at night. They may be found on the surface of ground which has been devoted for some years to lawn or sod purposes. The worms are usually much more numerous during the months of April, May, and June than at any other time, although they may be easily brought to the surface at any season of the year, except winter, by thoroughly sprinkling the soil in the early evening. If food is provided for the worms in the box, they may be kept almost indefinitely in such container without changing the soil. They have been raised successfully by feeding ordinary molasses spread on one side of a gunny sack, which is then laid on the surface of the ground with the sticky side downward, and the back of the bag then sprinkled with water. Powdered bread crumbs and crumbled hard boiled eggs have also been used as food.

Fresh Water Crawfish and Shrimp, Keeping Them Alive for Bait.—These crustaceans can be kept alive in tanks, small pools, or wooden boxes which are well supplied with running water. The best food for them is fresh meat fed in small pieces, but great care should be taken not to leave old and spoiled meat in the water for any length of time, as this will soon prove fatal. The boxes or other containers should not be overcrowded and should be cleaned often and the dead crawfish or shrimp thrown out, as they decay rapidly and will soon cause the death of the healthy ones. The same general treatment is used if the crustaceans are to be kept in closed tanks or aquaria.

Hellgramites.—These are the larval form of the dobson fly. They are found under stones in swift streams and are an excellent bait for bass. Hellgramites can be kept alive for a considerable time in floating bait boxes or in wet grass.

Glow Worms .- The term glow worm is

applied to the wingless female beetles of the family Lampyridae. They are nocturnal in habit and feed upon smaller insects and worms. They can be kept alive in loose, damp earth, covered with moist grass and kept in a cool place.

Preserving Minnows for Bait.—Take 1 part of formalin to 29 parts of water, place the minnows in this solution in a tightly closed jar or bottle and keep in the dark until they are to be used. In this way they will retain their colors and silvery hues better than if in the light.

When about to use the bait, soak it in fresh water to remove the formalin. A few drops of oil of rhodium may then be placed on the minnow to disguise the pungent odor of formalin that may remain in the fish after soaking. The oil of rhodium is said to be attractive to fish but be that as it may it does not repel them as the formalin is likely to do.

Dough Balls.—A tough paste may be made of moistened bean, wheat, or other flour, thoroughly mixed with a little sugar, or preferably honey. To give the paste a greater tenacity, cotton batting or wool should be stirred in. Ground or mashed white meat, such as veal or pork, or any bleached meat may be added, but this bait must be fresh and kept untainted. Dough balls may be made also by boiling rye flour to a consistency of paste, then sprinkling with corn flour and rolling into a "ball."

List of Common Fresh Water Game Fishes with General Mention of Some Baits Used in Their Capture

Bowfin, Dogfish, Grindle

Frogs, minnows, pieces of fish, etc. Blue Cat, Chuckle-Headed Cat, Fulton

Minnows, shiners, worms, crawfish, pieces of fish, meat, liver. Spotted Catfish, Channel Cat, Fiddler

Spotted Catfish, Channel Cat, Fiddler Shiners, worms, meat, liver, dough balls.

Common Bullhead, Brown Bullhead, Speckled Bullhead

Minnows, worms, frogs, grasshoppers, pieces of fish (chub, perch, sunfish), salt mackerel, salt pork, meat, liver.

Mud Cat, Yellow Cat, Goujon, Morgan Cat

Crawfishes, fresh hickory shad, other fish baits.

Buffalo Fish

Worms, insects. Carp Sucker

Worms, insects. Sucker

Earthworms, bits of crawfish.

Redhorse

Worms, insects.

Chub

Pieces of fish, insects, grasshoppers, worms.

Squawfish

Worms, minnows, shiners.

German Carp

For angling, various baits have been recommended. Worms, grubs, grasshoppers, and pieces of fresh meat have been used successfully, but the most highly recommended baits are composite pastes. Pellets of partly boiled potatoes are said to be good bait, as well as dough balls or corn kernels wrapped in mosquito bar.

American Eel, Fresh Water Eel Earthworms, shiners, grasshoppers, etc.

Mooneye

Minnows, worms, insects.

Common Whitefish

Worms, insect larvae, may flies, shrimp, pieces of fish, minnows.

Rocky Mountain Whitefish Worms, insects, fresh meat.

Salmon, Sea Salmon, Eastern Salmon Worms, smelt, shiners, pork rind.

Landlocked Salmon Smelts, shiners, worms.

Black Spotted Trout, Cut Throat Trou Worms, grasshoppers, insects, minnows, pieces of meat.

Steel Head Trout

Shiners, worms, insects, grasshoppers.

Rainbow Trout

Worms, grasshoppers, insects, shiners.

Brown Trout

Worms, various insects, grasshoppers, crickets, shiners, minnows, pieces of fish, horse meat.

Loch Leven Trout

Worms, various insects.

Chinook Salmon

Smelts, shiners.

Brook Trout

Earthworms or "barnyard hackle," grasshoppers, grubs, crickets, beetles, bumblebees, caterpillars, mill fly, caddis fly larvae, may fly, newts, mice, or bits of animal, flesh. A capital bait is the beautifully tinted anal fin of a trout, which in water with some current waves wabbles and flutters in a most seductive manner on the hook.

White Trout, Golden Trout

Worms, pieces of fish, smelts, minnows, shiners.

Dolly Varden Trout

Worms, minnows, shiners, insects.

Lake Trout

Minnows, shiners, pieces of fish (Whitefish), ciscoes.

Gravling

Gaddis fly larvae, "rock worm," earthworms, grubs, crickets, grass-hoppers, natural flies, or small bits of fat meat.

Smelt.

Pieces of smelt, shiners, minnows, worms, shrimp.

Common Pike. Pickerel

Frogs, shiners, minnows, white chub, pork rind, fish belly, 3-4 in. piece pickerel stomach, perch belly.

Muskellunge Small fishes, suckers, shiners, ciscoes, grasshoppers, frogs.

White Crappie

Worms, minnows, shiners. Black Crappie, Calico Bass Minnows, worms, small shiners.

Rock Bass, Redeye, Goggle-Eye Small minnows, white grubs, earthworms, grasshoppers, crickets, small crawfish, yellow perch, fresh water mussel, frogs.

Warmouth Bass (See Rock Bass.)

Red Robin, Long Eared Sunfish Worms, insects, minnows.

Bluegill, Blue Sunfish

Worms, insects, insect larvae, shrimps, small crawfish, pieces of fresh water mussel.

Green Sunfish, Blue Spotted Sunfish Worms, insects, insect larvae.

Pumpkinseed

Worms, insects, pieces of crawfish, pieces of meat.

Shell Cracker

Worms, insects, small crawfish, pieces of fish.

Black Bass

The best natural bait is the minnow, a shiner, chub, or the young of almost any fish, which is well adapted for either casting, trolling, or still fishing. In waters where it abounds, the crawfish is a good bait, especially the shedders or soft craws, to be used only for still fishing. The hellgramite, the larva of the corydalis fly, in its native waters, is also successful for still fishing. small frog is capital bait n weedy waters, where it is usually cast overhead with a very short and stiff rod. Grasshoppers and crickets are sometimes employed with a fly-rod, in lieu of artificial flies, with good results. The salt water shrimp, where it is available, near the coasts, is also a good bait for still fishing. Cut bait is also sometimes useful. It should be remembered that all baits of whatever kind, should be kept in motion. A dead minnow answers as well as a live one for casting or trolling, but should be alive for still With worms. fishing. crawfish. shrimps or hellgramites, a float should be employed to keep them from touching the bottom. In casting the minnow it should be hooked through the lips, and reeled in slowly after each cast to imitate the motions of a live one as much as possible.

Large Mouth Black Bass, Oswego Bass Live minnows and other live baits, such as grasshoppers, frogs, hellgramites, efts, worms.

Small Mouth Black Bass

Shiners, chub, small yellow perch with dorsal fin cut off, mad-tom, stone catfish, floor of mouth of pickeral cut like a fish, belly of bowfin, crawfish, hellgramites, crickets, efts, newts, small frogs, worms.

Wall Eyed Pike, Pike Perch, Jack

Salmon

Live minnows, as fallfish or dace, corporal, roach, redfin, gudgeon, brook chub, piece of fish, worms.

Yellow Perch, Ringed Perch, American Perch

Worms, minnows, crickets, grasshoppers and other insects, small fishes, small frogs, crawfish, pieces of fish. Striped Bass, Rockfish

Shiners, minnows, pieces of fish.

White Bass

Live minnows, grubs, earthworms.

Yellow Bass

Minnows (live bait), worms.

White Perch

Worms, grasshoppers, insects, minnows.

Fresh Water Drum, Croaker, Sheepshead, White Perch

Crawfish, pieces of fish, mollusks. Burbot, Ling, Eel Pout, Cusk

Yellow perch, sunfish, lamprey, crawfishes, pieces of fish, smelts.

Cut Flower Vitalizer U. S. Patent 1,978,201

Eight ounces of sugar or saccharin, 2 oz. of kaolin, 1 oz. of yeast, ½ oz. of charcoal, 1 cc. of oil of pine and ½ oz. of lime. The foregoing makes up a composition weighing about 12 oz. and this may be dissolved in a suitable amount of water. It has been found in practice that this diluted solution shows benefit to all cut flowers.

The benefit is so decisive that increased turgidity and intensified color in the tissue of leaf and petal are visible to the eye usually within 30 minutes after the flower stem is immersed in the diluted solution. This increased turgidity and intensified color is retained by the flower whether under average room temperature of 70° F. or in refrigerated temperatures of $40\text{--}50^{\circ}$ F., although a cooler temperature, as when untreated, prolongs the life of the cut flower.

The treated cut flower under observation slowly continues its development, retaining a healthy and nourished appearance, to eventually produce seed as large and apparently vital as it would upon the parent plant which had been un-

usually well cared for.

Furthermore, a flower cut in the bud develops normally when treated in this solution; for instance the chrysanthemum cut when the bud first shows color will develop into a flower equal in every respect to its companions left uncut on the greenhouse bench.

Again the treated flower lasts much longer after being removed from water, in treatment such as florists must subject

flowers to in funeral pieces.

Preserving Foliage

A method of preserving foliage consists in placing the leaves in a solution of glycerin 1 part, water 9 parts. The leaves are then removed from this solution, dried between blotting paper and pressed.

Gas Mask for Sulphur Dioxide

Flannel nose-bag masks, 7 in. by 8 in. and held over the face by rubber bands, are used as a protection against sulphur dioxide gas. Masks are soaked in the following solution:

TIO WINE DOLUMINA.	
Distilled Water	1000 cc.
Glycerin	250 cc.
Soda Ash	200 g.

Masks are worn while wet with the solution.

Gas-Producing Material for Inflating Hollow Rubber Articles British Patent 416,591

A mixture of sodium nitrite 56.5, ammonium chloride 43.5, and ammonium carbonate 10 parts is inserted into hollow rubber articles prior to vulcanization; on heating carbon dioxide, ammonia and nitrogen are evolved which ex-

pand the article up to the mold during vulcanization.

Manufacture of Luminescent Materials British Patent 414,905

A 2 to 1 mixture of zinc oxide (or magnesium oxide) and germanium dioxide is moistened with dilute aqueous manganese chloride and sintered at 1000° F. to produce zinc (or magnesium) germanate, which fluoresces bright greenishyellow (or orange-scarlet) under excitation with cathode rays.

Match-Striking Surfaces British Patent 411,688

An ignition surface, suitable for self-lighting cigarettes, etc., comprises a mixture of amorphous phosphorus and a cellulose derivative binder of the character of cellulose acetate. A mixture of 4 g. of amorphous phosphorus in 25 cc. of a 5% acetone solution of cellulose acetate is spread as a film on a suitable surface. Other solvents, e.g., ethyl acetate, may be used.

Microscope Slide Cleaner

Xylol	1	fl.	oz.
n-Butyl Alcohol	1	fl.	oz.
Alcohol, Anhydrous	2	fl.	oz.
Water	1	fl.	oz.

Sterile Modelling Clay U. S. Patent 1,979,016

Seventy grains of chlorthymol for every 100 lb. of manufactured modeling clay are sufficient to render the same sterile and to preserve its hygienic condition for long periods of time. The finished product may be packed in airtight containers for shipment and storage to prevent possible oxidation of its ingredients.

Preserving Fluid for Museum

···F··		
Formaldehyde	12-25	oz.
Glycerin	10	oz.
Potassium Nitrite	e 0.1	oz.
Water	to make 100	OZ.

Removing Formaldehyde Odor from Museum Specimens

Wash with water and submerge for 1/2 hour in:

Urea	5 oz.
Ammonium Phosphate	1 oz.
Water	94 oz.
If the specimen is to b	e replaced in

formaldehyde it should be washed free of urea.

Colored Neon Lights U. S. Patent 1,951,006

A mixture of approximately 10% of argon with 90% of neon emits a lavender The proportions of the colored light. gases may vary widely, the colors and shades changing with the different com-It is preferable to employ from 5 to 25% of argon, the balance being principally neon. The addition of carbon dioxide to the mixture of neon and argon, for example, results in a white or substantially colorless light. Therefore, introduce a substance such as calcium or magnesium carbonate, which is capable of releasing carbon dioxide to the tube containing rare gases such as neon and argon. When the tube is energized, carbon dioxide is released, and produces the white or substantially colorless light until the modifying agent is exhausted. Such tubes have been operated for more than 700 hours without change of the light emitted.

In introducing the modifying agent to the tube, several methods may be employed: The agent may be supported inside the electrode; it may be attached to the electrode; it may be coated on the wall of the tube or electrode chamber; or it may be simply deposited in the electrode chamber or in the path of the discharge through the tube.

Other modifying agents may be used, for example, a suitable hydride such as magnesium hydride can be used to maintain a trace of hydrogen in the tube in admixture with the gases therein to effect a desired change in the color of the light emitted when the tube is energized.

Electrode, Neon U. S. Patent 1,926,336

The electrode comprises a compressed cylinder of an intimate mixture of tantalum carbide (88%) and cesium chloride, rubidium chloride and lithium chloride (12%).

Oxalic Acid from Corncobs

Corncobs 100 lb. Nitric Acid (95%) 3 lb.

Heat until dissolution is complete; cool and add:

Nitric Acid (50-55%) 3 lb. Vandium Pentoxide 0.1 lb. Allow to stand for 2 or 3 days; filter and evaporate the filtrate to obtain crude oxalic acid which may be purified by recrystallization.

Radiator Corrosion Inhibitor U. S. Patent, 1,992,689

For preventing corrosion in motor radiators containing alcohol and water the following formula is used:

Triethanolamine Triethanolamine Phosphate	0.33	oz.
phate	1.50	oz.
b. Triethanolamine Lard Oil	0.75	oz.
Lard Oil	0.75	oz.

Mix ingredients of b and stir into a. The above is used per 100 parts of alcohol.

Scale Preventing Mixture

Formula No. 1 French Patent 776,235

A mixture of formic acid 100 and digallic acid 6 parts is used.

No. 2 French Patent 776,234

A mixture of digallic acid 100, and trisodium phosphate 60 parts, is used to prevent scale in motor car radiators.

Non-Corrosive Chlorinated Solvents U. S. Patent 1,966,881

Five-tenths to 2% of pinene is added to prevent corrosion.

Tellurium Alloy Rectifier U. S. Patent 1,961,825

The rectifier consists of plates of magnesium and an alloy of:

Tellurium	97.5	oz.
Copper	2	oz.
Silver	2.5	OZ.
Sodium	0.5	oz.

which are welded together by passing a current from one to the other with a film of water between them.

Aluminum Reflector Etching U. S. Patent 1,999,042

Using hydrofluoric acid and nitric acid the aluminum is first dipped into a solution of 1 part concentrated hydrofluoric acid in 19 parts of water at a temperature of 50 to 60° C., until an etch of the desired depth is obtained. The surface is washed and the article is im-

77 lb.

23 lb.

mersed for several seconds in a solution of nitric acid containing 1 volume of acid to 1 volume of water and held at room temperature. The aluminum is washed and dried and a clean, bright and uniformly etched surface is obtained. In the sodium hydroxide-sodium fluoride etching procedure a 5% sodium hydroxide solution in water containing about 4% sodium fluoride is used. The aluminum is immersed in this solution at a temperature of about 90° C. until the desired etch is obtained. It is then removed, washed, and treated with a 1:1 nitric acid solution, washed and dried as before. Again a very satisfactory clean, bright, and uniformly etched surface is obtained. It should be noted that the presence of copper in the aluminum causes the metal to turn gray to black on immersion in the hydrofluoric acid or the sodium hydroxide solutions. black coloration, due to copper, is removed by immersion in the nitric acid. However, the nitric acid does not remove the gray film due to graphitic silicon, if it is present, and this must either be removed by rubbing or prevented from forming.

The effect of the presence of a sufficient amount of copper in aluminum on its etching properties is pointed out specifically by the following examples: A sample of a commercial grade of aluminum containing about 1% of impurities, including 0.6% iron, 0.3% silicon, and 0.01% copper, when etched with hydrofluoric and nitric acids as above described, produces a surface which is irregularly etched, having a streaked appearance. On the other hand, a sample of aluminum containing 0.6% iron, 0.08% copper, and 0.18% silicon as impurities, when etched in a similar manner, produces a very satisfactory uni-

form reflecting surface.

Brine for Refrigeration U. S. Patent 1,969,124

A eutectic solution for refrigerating purposes comprises barium chloride 19, potassium chloride 18 and sodium chloride 4 oz. per gallon of water.

Refrigerator Deodorant

Fill a small muslin bag with a good quality of granular activated carbon. The muslin bag may then be placed in the rear of a lower portion of the ice box and will absorb strong odors which tend to collect.

After six months use, the device may

be reactivated by placing in the oven at 350° F. for about ½ hour.

Increasing Resistance of Magnesium Oxide

U. S. Patent 2,012,897

A process for increasing the electrical resistivity of fused magnesium oxide comprises heating magnesium oxide in an oxidizing atmosphere for approximately 6 hours at a temperature of approximately 2000–2300° F.

Salt Denaturant

Two per cent of wormwood powder is added to salt for industrial use.

Soot Destroyer Canadian Patent 347,077

Lead Oxide Salt

The above may be diluted with charcoal or sawdust.

Stop Leak Composition U. S. Patent 1,988,764

A stop leak composition for water circulating systems, comprising as chief ingredients about 4 g. of paper pulp, 5 g. of sifted flax seed, 200 cc. of water, and a small percentage of a preservative.

Temperature Sensitive Compounds

The following color changes induced by temperature changes find applications in many fields:

- 1. Copper Ferrocyanide.—Is manogany brown at room temperature, becomes brown-black on heating, returns to original color on cooling.
- 2. Arsenic Bisulphide.—Orange red at room temperatures, changes progressively to dark red and then brown at higher temperatures, returns to original color on cooling.
- 3. Lead Iodide. Original orange changes to dark orange on heating.
- 4. Mercury Subsulphide.—Original yellow changes on heating to orange yellow, then orange, then red.
- 5. Lead Chromate.—Same changes as for mercury subsulphide.
- 6. Tin Subsulphide.—Original brown color (or orange yellow) changes to dark red, then nearly black, on heating. These changes are very temperature sensitive.

- 7. Silver Subiodide.—Green yellow at ordinary temperatures changes to orange when heated.
- 8. Mercury Subiodide.—Original yellowish green changes on heating to orange, red, and brownish red.
- 9. Weak Copper Bromide.—Original lemon-yellow turns to brown when heated, returning to original color when cooled.
- 10. Cobalt Chloride.—Is invisible at ordinary temperatures but becomes blue when heated.
- 11. Mercuric Oxide.—Red at ordinary temperatures, darkens on heating, becomes black eventually.

Thermionic Cathode U. S. Patent 1,961,122

The filament consists of an alloy of: Nickel 90 oz. Iron 7.5 oz. Titanium 2.5 oz.

Coated with barium oxide.

Protecting Carbide

Carbide will keep indefinitely if sprinkled uniformly with kerosene.

Tooth Desensitizer (Hartman)

Ether 2 oz. Alcohol 1 oz. Thymol 1¼ oz.

Keep in a brown bottle, tightly stoppered.

Apply inside of tooth by means of a dab of absorbent cotton on a tooth pick. The cavity in which it is applied should be dry to insure lengthy desensitization. Contact should be for 1 to 1½ minutes. The cotton is then removed and the cavity is dried with a blast of hot air.

Denicotinized Cigarettes

Activated charcoal and silica gel is used in individual cigarettes for the absorption of nicotine. Charcoal (0.2 g.) or silica gel (0.1 g.) is an efficient denicotinizer.

Denicotinizing Tobacco U. S. Patent 2,000,855

A method of denicotinizing tobacco comprises the steps of: wetting tobacco containing the usual bacteria, disposing the wetted tobacco loosely in layers and allowing the latter to stand with access of air thereto to produce fermentation of the tobacco, continuously adding acid

to the extent necessary to neutralize the amino bases resulting from the fermentation, and drying the tobacco.

Treating Tobacco for Smoking U. S. Patent 1,972,718

There is added to tobacco about 2% of an alkaline hydrated aluminum silicate which upon the smoking of the tobacco is capable of taking up gases and tarry compounds produced by the combustion.

Water-Softening Compound U. S. Patent 1,952,408

A cake for domestic use, formed by pressure when moist, comprises sodium carbonate 62.5, sodium phosphate 30.0, calcium chloride 5.0, and sodium chloride 2.5%.

Base-Exchange Materials for Water Softening

British Patent 434,663

Raw clay is treated with concentrated hydrochloric or sulphuric acid, the supernatant acid removed, and the clay baked at 550-600° F. for 1 hour. The product is treated with 10% aqueous sodium silicate, then with 2% aqueous sodium aluminate at 100° F., and finally with 5% aqueous sodium chloride to increase the base exchange power.

Water Testing Indicator British Patent 414,836

The dipping rod is coated with a paste made from chalk (16), glycerin (12), a saturated solution of rosin in turpentine (1), and methylene blue dissolved in methylated spirit (1); contact with water lightens the color.

Windshield Anti-Fog Compound Formula No. 1

Windshields may be kept clear of fog, by occasionally wiping them with a cloth prepared by boiling it 10 minutes in a solution of:

Water 5 qt.
Glycerin 1 oz.
Sodium Oleate 1 oz.

Boil together 5 minutes before immersing cloth.

No. 2

Glycerin 10 oz.
Glycol Boriborate 4 oz.
Sulphonated Castor Oil 10 drops

TABLES

Weights and Measures Troy Weight

24 grains = 1 pwt. 20 pwts. = 1 ounce 12 ounces = 1 pound

Apothecaries' Weight

20 grains = 1 scruple 3 scruples = 1 dram8 drams = 1 ounce12 ounces = 1 pound

The ounce and pound are the same as in Troy Weight.

Avoirdupois Weight

2711/32 grains = 1 dram 16 drams = 1 ounce 16 ounces = 1 pound2000 lbs. = 1 short ton 2240 lbs. = 1 long ton

Dry Measure

2 pints = 1 quart 8 quarts = 1 peck 4 pecks = 1 bushel 36 bushels = 1 chaldron

Liquid Measure

4 gills = 1 pint2 pints = 1 quart4 quarts = 1 gallon $31\frac{1}{2}$ gals. = 1 barrel 2 barrels = 1 hogshead 1 teaspoonful = 1/6 oz. 1 tablespoonful $= \frac{1}{2}$ oz. 16 fluid oz. = 1 pint

Circular Measure

60 seconds = 1 minute 60 minutes = 1 degree 360 degrees = 1 circle

Long Measure

12 inches = 1 foot3 feet = 1 yard $5\frac{1}{2}$ yards = 1 rod 5280 feet = 1 stat. mile 320 rods = 1 stat. mile Square Measure

144 sq. in. = 1 sq. ft.9 sq. ft. = 1 sq. yard $30\frac{1}{4}$ sq. yds. = 1 sq. rod 43,560 sq. ft. = 1 acre40 sq. rods = 1 rood4 roods = 1 acre640 acres = 1 sq. mile

Metric Equivalents

Length

1 inch = 2.54 centimeters 1 foot = 0.305 meter1 yard = 0.914 meter1 mile = 1.609 kilometers1 centimeter = 0.394 in. 1 meter = 3.281 ft.1 meter = 1.094 yd.1 kilometer = 0.621 mile

Capacity

1 U.S. fluid oz. = 29.573 milliliters 1 U. S. liquid qt. = 0.946 liter 1 U. S. dry qt. = 1.101 liters 1 U. S. gallon = 3.785 liters 1 U. S. bushel = 0.3524 hectoliter 1 cu. in. = 16.4 cu. centimeters 1 milliliter = 0.034 U.S. fluid ounce 1 liter = 1.057 U. S. liquid qt. 1 liter = 0.908 U. S. dry qt.1 liter = 0.264 U. S. gallon1 hectoliter = 2.838 U. S. bu. 1 cu. centimeter = .061 cu. in. 1 liter = 1000 milliliters or 100 cu. c.

Weight

1 apoth. scruple = 1.296 grams 1 av. oz. = 28.350 grams1 troy oz. = 31.103 grams 1 av. lb. = 0.454 kilogram 1 troy lb. = 0.373 kilogram1 gram = 15.432 grains1 gram = 0.772 apoth. scruple1 gram = 0.035 av. oz.1 gram = 0.032 troy oz.1 kilogram = 2.205 av. lbs. 1 kilogram = 2.679 troy lbs.

1 grain = 0.065 gram

Annrovimata nH Values	Reets	4.9-5.5
Approximate pH Values		3.2-3.6
The following tables give approximate		
pH values for a number of substances		5.0-6.0
such as acids, bases, foods, biological		6.1-6.4
fluids, etc. All values are rounded off to		5.2 - 5.4
the nearest tenth and are based on meas-	Carrots	4.9 - 5.3
urements made at 25° C.	Cheese	4.8 - 6.4
	Cherries	3.2 - 4.0
pH Values of Acids	Cider	2.9 - 3.3
Hydrochloric, N 0.1	Corn	6.0 - 6.5
Hydrochloric, N 0.1	Crackers	6.5-8.5
Hydrochloric, 0.1N 1.1 Hydrochloric, 0.01N 2.0	Dates	6.2-6.4
Hydrochloric, 0.01N 2.0		
Sulphuric, N 0.3	Eggs, fresh white	7.6-8.0
Sulphuric, N 0.3 Sulphuric, 0.1N 1.2 Sulphuric, 0.01N 2.1	Flour, wheat	5.5-6.5
Sulphuric, 0.01N 2.1		2.8 - 3.0
Orthophosphoric, 0.1N 1.5 Sulphurous, 0.1N 1.5		3.0-3.3
Sulphurous. 0.1N	Grapes	3.5 - 4.5
Oxalic, 0.1N 1.6	Hominy (lye)	6.8 - 8.0
Oxalic, 0.1N 1.6 Tartaric, 0.1N 2.2	Jams, fruit	3.5 - 4.0
Malic, 0.1N 2.2	Jellies, fruit	2.8-3.4
	Lemons	2.2-2.4
Oldfic, U.IIN		
Formic, 0.1N	Limes	1.8-2.0
Lactic, 0.1N 2.4	Maple Syrup	6.5-7.0
Lactic, 0.1N	Milk, cows	6.3-6.6
Acetic, 0.1N 2.9 Acetic, 0.01N 3.4	Olives	3.6–3.8
Acetic, 0.01N 3.4	Oranges	3.0 - 4.0
Benzoic, 0.1N 3.1	Oysters	6.1 - 6.6
Alum, 0.1N 3.2	Peaches	3.4 - 3.6
Cambania (setumated)	Pears	3.6 - 4.0
Carbonic (saturated) 3.8	Peas	5.8-6.4
Hydrogen Sulphide, 0.1N 4.1	Pickles, dill	3.2-3.6
Arsenious (saturated) 5.0	Dickles, dill	
Hydrocyanic, 0.1N 5.1	Pickles, sour	3.0-3.4
Boric, 0.1N 5.2	Pimento	4.6-5.2
	Plums	2.8 - 3.0
pH Values of Bases	Potatoes	5.6 - 6.0
A 7 7 7 17 37	Pumpkin	4.8 - 5.2
Sodium Hydroxide, N 14.0	Raspberries	3.2 - 3.6
Sodium Hydroxide, N 14.0 Sodium Hydroxide, 0.1N 13.0 Sodium Hydroxide, 0.01N 12.0	Rhubarb	3.1 - 3.2
Sodium Hydroxide, 0.01N 12.0	Salmon	6.1-6.3
Potassium Hydroxide, N 14.0	Sauerkraut	3.4-3.6
Potassium Hydroxide, 0.1N 13.0	Shrimp	6.8-7.0
Potassium Hydroxide, 0.01N 12.0		
	Soft Drinks	2.0-4.0
Lime (saturated) 12.4 Sodium Metasilicate, 0.1N 12.6	Spinach	5.1 - 5.7
Trisodium Phosphate, 0.1N 12.0	Squash	5.0 - 5.4
	Strawberries	3.0-3.5
Sodium Carbonate, 0.1N 11.6	Sweet Potatoes	5.3 - 5.6
Ammonia, N 11.6	Tomatoes	4.0 - 4.4
Ammonia, 0.1N 11.1 Ammonia, 0.01N 10.6	Tuna	5.9 - 6.1
Ammonia, 0.01N	Turnips	5.2 - 5.6
Potassium Cyanide, 0.1N 11.0	Vinegar	2.4-3.4
Magnesia (saturated) 10.5	Water, drinking	6.5-8.0
Sodium Sesquicarbonate, 0.1N 10.1	Wines	2.8-3.8
Ferrous Hydroxide (saturated) 9.5	Willes	4.0-5.0
Calcium Carbonate (saturated) 9.4	pH Values of Biologic Mater	rials
Borax, 0.1N 9.2		
	Blood, plasma, human	7.3 - 7.5
Sodium Bicarbonate, 0.1N 8.4	Spinal Fluid, human	7.3 - 7.5
pH Values of Foods	Blood, whole, dog	6.9 - 7.2
bir varies or rooms	Saliva, human	6.5 - 7.5
Apples 2.9–3.3	Gastric Contents, human	1.0-3.0
Apricots 3.6-4.0	Duodenal Contents, human	4.8-8.2
Asparagus 5.4–5.8	Feces, human	4.6-8.4
Bananas 4.5–4.7		
	Urine, human	4.8-8.4
Beans 5.0-6.0	Milk, human	6.6-7.6
Beers 4.0-5.0	Die, numan	6.8 - 7.0

TABLES

CONVERSION OF THERMOMETER READINGS

\mathbf{F}°	C°	F°	C°	F°	C°	F°	C°	F°	C°	F°	C°
-40 -38 -36 -34 -32	-40.00 -38.89 -37.78 -36.67 -35.56	33	$\begin{array}{c c} -1.11 \\ -0.56 \\ 0.00 \\ 0.56 \\ 1.11 \end{array}$	80 81 82 83 84	26.67 27.22 27.78 28.33 28.89	250 255 260 265 270	121.11 123.89 126.67 129.44 132.22	500 505 510 515 520	260.00 262.78 265.56 268.33 271.11	900 910 920 930 940	482.22 487.78 493.33 498.89 504.44
-30 -28 -26 -24 -22	$\begin{array}{r} -34.44 \\ -33.33 \\ -32.22 \\ -31.11 \\ -30.00 \end{array}$	35 36 37 38 39	1.67 2.22 2.78 3.33 3.89	85 86 87 88 89	29.44 30.00 30.56 31.11 31.67	285 290	135.00 137.78 140.55 143.33 146.11	525 530 535 540 545	273.89 276.67 279.44 282.22 285.00	960 970 980	510.00 515.56 521.11 526.67 532.22
-20 -18 -16 -14 -12	-28.89 -27.78 -26.67 -25.56 -24.44	40 41 42 43 44	4.44 5.00 5.56 6.11 6.67	90 91 92 93 94	32.22 32.78 33.33 33.89 39.44	315	148.89 151.67 154.44 157.22 160.00	550 555 560 565 570	287.78 290.55 293.33 296.11 298.89	1050 1100 1150	
$ \begin{array}{r} -10 \\ -8 \\ -6 \\ -4 \\ -2 \end{array} $	-23.33 -22.22 -21.11 -20.00 -18.89	45 46 47 48 49	7.22 7.78 8.33 8.89 9.44	95 96 97 98 99	35.00 35.56 36.11 36.67 37.22	330 335 340	162.78 165.56 168.33 171.11 173.89	580 585 590	301.67 304.44 307.22 310.00 312.78	1300 1350 1400	732.22 760.00
0 1 2 3 4	-17.78 -17.22 -16.67 -16.11 -15.56	53	10.00 10.56 11.11 11.67 12.22	100 105 110 115 120	37.78 40.55 43.33 46.11 48.89	355 360 365	176.67 179.44 182.22 185.00 187.78	610 620 630	315.56 321.11 326.67 332.22 337.78	1550 1600 1650	843.33 871.11 898.89
5 6 7 8 9	-15.00 -14.44 -13.89 -13.33 -12.78	56 57 58	12.78 13.33 13.89 14.44 15.00	125 130 135 140 145	51.67 54.44 57.22 60.00 62.78	380 385 390	190.55 193.33 196.11 198.89 201.67	660 670 680	343.33 348.89 354.44 360.00 365.56	1800 1 1850 1 1900	
10 11 12 13 14	-12.22 -11.67 -11.11 -10.56 -10.00	61 62 63	15.56 16.11 16.67 17.22 17.78	155 160	65.56 68.33 71.11 73.89 76.67	410	204.44 207.22 210.00 212.78 215.56	710 720 730	371.11 376.67 382.22 387.78 393.33	2050 2 2100 3 2150	1093.33 1121.11 1148.89 1176.67 1204.44
15 16 17 18 19	- 9.44 - 8.89 - 8.33 - 7.78 - 7.22	66 67	18.33 18.89 19.44 20.00 20.56		79.44 82.22 85.00 87.78 90.55	430 435 440	218.33 221.11 223.89 226.67 229.44	760 770 780	398.89 404.44 410.00 415.56 421.11	2300 2350 2400	1232.22 1260.00 1287.78 1315.56 1343.33
20 21 22 23 24	- 6.67 - 6.11 - 5.56 - 5.00 - 4.44	71 72 73	21.11 21.67 22.22 22.78 23.33	215	93.33 96.11 98.89 101.67 104.44	455 460 465	232.22 235.00 237.78 240.55 243.33	810 820 830	426.67 432.22 437.78 443.33 448.89	2 2550 3 2600 3 2650	1371.11 1398.89 1426.67 1454.44 1482.22
25 26 27 28 29	- 3.89 - 3.33 - 2.78 - 2.22 - 1.67	76 77 78	23.89 24.44 25.00 25.56 26.11	230 235 240	107.22 110.00 112.78 115.56 118.33	485	246.11 248.89 251.67 254.44 257.22	860 870 880	454 . 44 460 . 00 465 . 50 471 . 1 476 . 67	2800 3 2850 1 2900	1510.00 1537.78 1565.56 1593.33 1621.11

ALCOHOL PROOF AND PERCENTAGE TABLE

U.S. Proof at 60° F.	Per cent Alcohol by Volume at 60° F.	Per cent Alcohol by Weight	U.S. Proof at 60° F.	Per cent Alcohol by Volume at 60° F.	Per cent Alcohol by Weight
0	0.0	0.00	58	29.0	23.82
1 2 3 4 5 6 7 8	0.5	0.00	59	29.5	0.4.05
3	1.0 1.5	0.80	60 61	30.0 30.5	24.67
4	2.0	1.59	62	31.0	25.52
5	$\widetilde{2.5}$		63	31.5	
6	3.0	2.39	64	32.0	26.38
7	3.5	0.70	65	32.5	
9	4.0 4.5	3.19	66 67	33.0 33.5	27.24
10	5.0	4.00	68	34.0	28.10
11	5.5		69	34.5	-
12	6.0	4.80	70	35. 0	28.97
$\frac{13}{14}$	6.5	F 07	71	35.5	20.04
15	7.0 7.5	5.61	$\begin{array}{c} 72 \\ 73 \end{array}$	36.0 36.5	29.84
16	8.0	6.42	74 74	37.0	30.72
17	8.5		75	37.5	
18	9.0	7.23	76	38.0	31.60
19 20	9.5	0.05	77	38.5	00.40
$\frac{20}{21}$	$10.0 \\ 10.5$	8.05	78 79	39.0 39.5	32.48
22	11.0	8.86	80	40.0	33.36
23	11.5		81	40.5	
24	12.0	9.68	82	41.0	34.25
$\begin{array}{c} 25 \\ 26 \end{array}$	$12.5 \\ 13.0$	10.50	83	41.5	05.15
27 27	13.5	10.50	84 85	$42.0 \\ 42.5$	35.15
28	14.0	11.32	86	43.0	36.05
29	14.5		87	43.5	
30	15.0	12.14	88	44.0	36.9 6
$\begin{array}{c} 31 \\ 32 \end{array}$	15.5 16.0	12.96	89 90	$\frac{44.5}{45.0}$	37.86
33	16.5	12.30	91	45.5	01.00
34	17.0	13.79	92	46.0	38.78
35	17.5		93	46.5	
36 37	18.0	14.61	94	47.0	39.7 0
38	18.5 19.0	15.44	95 96	47.5 48.0	40.62
39	19.5		97	48.5	40.02
40	20.0	16.27	98	49.0	41.55
41	20.5		99	49.5	
42 43	$\begin{array}{c} 21.0 \\ 21.5 \end{array}$	17.10	$\begin{array}{c} 100 \\ 101 \end{array}$	50.0 50.5	42.49
44	22.0	17.93	102	51.0	43.43
45	22.5		103	51.5	10.10
46	23.0	18.77	104	52.0	44.37
47	23.5	70.00	105	52.5	
48 49	$24.0 \\ 24.5$	19.60	106	53.0	45.33
50	25.0	20.44	$\begin{array}{c} 107 \\ 108 \end{array}$	53.5 54.0	46.28
51	25.5		109	54.5	±0.20
52	26.0	21.28	110	55.0	47.24
53	26.5	00.10	111	55.5	
54 55	$\begin{array}{c} 27.0 \\ 27.5 \end{array}$	22.13	$\begin{array}{c} 112 \\ 113 \end{array}$	56.0	48.21
56	28.0	22.97	113	56.5 57.0	49.19
57	28.5		115	57.5	10.10

U.S. Proof at 60° F.	Per cent Alcohol by Volume at 60° F.	Per cent Alcohol by Weight	U.S. Proof at 60° F.	Per cent Alcohol by Volume at 60° F.	Per cent Alcohol by Weig!
116	58.0	50.17	159	79.5	
117	58.5	F1 15	160	80.0	73.53
118	59.0 59.5	51.15	$161 \\ 162$	80.5 81.0	74.69
$\frac{119}{120}$	60.0	52.15	163	81.5	74.09
121	60.5	02.10	164	82.0	75.86
$\frac{121}{122}$	61.0	53.15	165	82.5	70.00
123	61.5		166	83.0	77.04
124	62.0	54.15	167	83.5	
125	62.5		168	84.0	78.23
126	63.0	55.1 6	169	84.5	
127	63.5	***************************************	170	85.0	79.44
128	64.0	56.18	171	85.5	
129	64.5		172	86.0	80.62
130	65.0	57.21	173	86.5	
131	65.5		174	87.0	81.90
132	66.0	58.24	175	87.5	
133	66.5		176	88.0	83.14
134	67.0	59.28	177	88.5	04.41
135	67.5	40.00	178	89.0	84.41
136	68.0	60.32	1.79	89.5	05.00
137	68.5	01.00	180	90.0	85.69
138	69.0	61.38	181	90.5	86.99
139	69.5	60.44	182	91.0	50.99
140	70.0	62.44	183	$91.5 \\ 92.0$	88.31
141	70.5	63.51	184	92.0 92.5	90.01
142	$71.0 \\ 71.5$	09.91	185 186	92.5 93.0	89.65
143	$\begin{array}{c} 71.5 \\ 72.0 \end{array}$	64.59	187	93.5	
$\frac{144}{145}$	72.5	04.05	188	94.0	91.02
$\frac{145}{146}$	73.0	65.67	189	94.5	
147	73.5		190	95.0	92.42
148	74.0	66.77	191	95.5	
149	74.5		192	96.0	93.85
150	75.0	67.87	193	96.5	
151	75.5		194	97.0	95.32
152	76.0	68.92	195	97.5	-
153	76.5		196	98.0	96.82
154	77.0	70.10	197	98.5	
155	77.5		198	99.0	98.38
156	78.0	71.23	199	99.5	
157	78.5	·····	200	100.0	100.00
158	79.0	72.38	1		

Buffer Systems

The following table gives some common buffer systems and the approximate pH of maximum buffer capacity. The zone of effective buffer action will vary with concentration but the general average will be \pm 1.0 pH from the value given, for concentrations approximately 0.1 molar.

Glycocoll - Sodium Chloride - Hydro-	4
chloric Acid	2.0
Potassium Acid Phthalate-Hydro-	
chloric Acid	2.8
Primary Potassium Citrate	3.7
Acetic Acid-Sodium Acetate	4.6

Potassium Acid Phthalate-Sodium	
Hydroxide	5.0
Secondary Sodium Citrate	5.0
Carbonic Acid-Bicarbonate	6.5
Primary Phosphate-Secondary Phos-	
phate	6.8
Primary Phosphate-Sodium Hydrox-	
ide	6.8
Boric Acid-Borax	8.5
Borax	9.2
Boric Acid-Sodium Hydroxide	9.2
Bicarbonate-Carbonate	10.2
Secondary Phosphate-Sodium Hy-	
droxide	11.5
Courtesy of W A Taulor & Com	mann

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Allg. Oes. v. Gettzeitung
Amer. Druggist
Amer. Dyestuff Reporter
Amer. Electrop. Society
Amer. Paint Jol.
Amer. Perfumer
Amer. Photography
Amer. Wool & Cotton Reporter
Analyst
Anal. Fis. Quim.
Ault & Wiborg Varnish Wks. Handbook

Baker's Helper
Bakers Review
Baker's Weekly
Better Enameling
Bottler & Packer
Boyce Thompson Inst.
Brewers' Tech. Review
Brick & Clay Record
Br. Jol. Dent. Science
Brit. Jol. of Photography
Brit. Medical Jol.
Bull. Imp. Hyg. Lab.
Bulletin of Imperial Institute
Bull. Soc. Franc. Phot.

Camera
Camera (Luzern)
Canner
Cement & Cement Mfr.
Chemical Abstracts
Chemical Analyst
Chemical Industries
Chemical Weekblad
Chem. Zent.
Chemist & Druggist
Combustion
Confectioner's Jol.
Cramer's Manual

Dairy World
Dansk. Tids. Farm
Dental Lab'y Review
Devt. Part. Zeitung
Drug & Cosmetic Industry
Druggists Circular
Drugs, Oils, & Paints

Eastman Kodak Co. Electric Journal

Farbe v. Lacks

Farben Zeitung
Farming S. Africa
Fein Mechanic v. Prazision
Fettchem, Umschan
Fils & Tissus
Focus
Food Manufacture
Fruit Products Jol.

Gelatin, Leim, Klebstoffe Glass Industry

Hawaiian Planters' Record Hide & Leather

Ice Cream Review
India Rubber World
Indian Lac Research Inst.
Industrial Chemist
Industrial Finishing
Int'l Tin Res. & Dev. Council

Jol. Amer. Dental Assn.
Jol. Amer. Medical Assn.
Jol. Chinese Chem. Soc.
Jol. Federation Curriers
Jol. Federation Light Leather Tanners
Jol. Ind. & Eng. Chemistry
J. Res. Nat. Bur. Standards
Jol. Rubber Industry
J. Russ. Rubber Ind.
Jol. Soc. Leather Trades
Jol. Soc. Rubber Ind. Japan

Keram Steklo Khimstroi Kozhevenna-Obuvnaya Prom. Kunstdunger, Und Leim

Lakokras, Ind. Leather Trades Review Les Mat. Grasses Lithographic Tech. Foundation

Malayan Agric. Jol.
Manufacturing Chemist
Meat
Meat Merchandising
Melliand
Metal Industry
Metall und Erz
Metallurg
Metallurgist
Metalls & Alloys

Mich. Agric. Exp. Sta. Monatschr. Textil-Ind. Munic. Eng. San. Record

Nat'l Butter & Cheese Jol. Nat'l Provisioner Nickelsworth Nitrocellulose

Ober Flachen Tach. Oil & Color Trades Jol. Oil & Soap

Paper Trade Jol. Parfum Mod. Peinture, Pigments, Vernis Phar. Acta Helva Pharmaceutical Jol. Phot. Abstracts Phot. Ind. Phot. Korr. Photog. Kronik Phot. Rev. Photo Rundschau Physics Phytopathology Plater's Guide Book Portland Cement Assn. Power Practical Druggist Practical Everyday Chemistry Printing Industry
Prob. Edelmetalle
Process Engr. Mo.
Proc. World Petroleum Congress

Rayon & Mell. Tex. Monthly Refiner & Nat. Gas Mfr. Rev. Aluminum Rev. Amer. Electro Society Rock Products

Science Shoe and Leather Journal Soap Soap Gazette & Perfumer Sovet-News Sovet-Sakhar Spirits Synthetic & Applied Finishes

Textile Colorist Textile Mfr. Textile Recorder

U. S. Department of AgricultureU. S. Bureau of MinesU. S. Bureau of Standards

Veneers and Plywood

Z. Elektrochem. Zeit. Unters. Lebensm.

COMMON NAMES OF CHEMICAL PRODUCTS

Α

Acacia GumGum Arabic
Acetate of Lime
Acetic EtherEthyl Acetate
Acetin Acetate
Acetyl Salicylic Acid Glyceryl Monoacetate Acetyl pa Totrophloride Aspirin
Acetylane Tetrachloride
Acetylene Tetrachloride
Tadopo Landa e e e e e e e e e e e e e e e e e e e
Alcohol Fthyl Alcohol
Adminum Fotassium Siliphate Alim
Amount TT 7
Aniline
Animal Chargosl Aniline Oil
Animal Charcoal
Titmin A 1
And a Company of the company of the
AsphaltumMineral Pitch

В

Baking Soda	. Sodium Bicarbonate
	Posisson Calabata as a
Benzine Rlack Boy Gum	.Petroleum
Black Lead	· Accroides Gum
Bleaching Powder Blue Stone	. Calcium Hypochlorite
Blue Stone Blue Vitriol Boiled Oil	Connor Gul-1
Roiled Oil	Copper Sulphate
Boiled Oil Bone Black	.Boiled Linseed Oil
- Oldott Held	D
Doruk	D .
	C11
Burnt Sugar Coloring Butanol	Caramel Color
Butanol	Butyl Alcohol
Butyric Ether	Ethyl Butyrate

c

			Calcium Mercuro		Monobasic
Caoutcho	ac	 •••••	India Ru	us Chloride ibber	

Capsicum Carbolic Acid Carragheen Cartechu Caustic Potash Caustic Soda Ceresin Wax Calcium Carbonate China Clay China Wood Oil Chinese Wax Chloride of Lime Cinnabar Cinnabar Cinnabar Cinnabar Cinnabar Cinnabar Cinnabar Cinnabar Cinnabar Cinnabar Cinnabar Colloidal Clay Colloidal Clay Colloidal Clay Colloidal Clay Colloidal Clay Colloidal Clay Colloidal Clay Colloidal Clay Colloidal Clay Collogne Spirits Colloidal Clay Colopper Aceto Arsenite Corrosive Sublimate Coram of Tartar Cresol Crambin Spirats Colloid Crambin Colloide Corambin Colloide Coram of Tartar Cresol Crambin Colloide Calcium Hypochlor Coleptor Colophony Colopper Aceto Arsenite Colloidal Clay Colloidal Clay Corrosive Sublimate Corrosive Sublimate Corrosive Sublimate Corrosive Sublimate Coresol Cresylic Acid Crude Oil Calcium Cyanamide Calcium Cyanamide Calcium Cyanamide Calcium Cyanamide Calcium Cyanamide	ution'' re) te
D	

Dead Oil	Creosote Oil
Decalin	Decahydronaphthalene
Degras	Wool Grease
Dope	Pvroxvlin "solution"
Dutch Liquid	Ethylene Chloride

E

Earth, Infusorial	.Earth, Diatomaceous
Egg Oil	.Egg Yolk
Elaterite	. Mineral Rubber
Epsom Salts	. Magnesium Sulphate
Ether	
Ethvl Nitrite	.Nitrous Ether

F

Fir, Balsam Flaxseed Flea seed	Linseed
Fluorspar Fool's Gold Fool's	Calcium Fluoride Iron Pyrites
Formalin French Chalk	Tale
Fuchsine	Amyl Alcohol (fermentation amyl alcohol)

G

Galena Lead Sulphide Glance Pitch Manjak Glass, Water Sodium Silicate Glauber's Salt Sodium Sulphate Glycerin Glycerol Glycol Ethylene Glycol Graphite Plumbago Green Soap Soft Soap Green Vitriol Ferrous Sulphate Ground Nut Oil (Arachi's Oil) Peanut Oil Gum Lac Shellac Gun Cotton Nitro-Cellulose Gypsum Calcium Sulphate
H
Heavy Spar Barium Sulphate Hematite Iron Oxide Hexamine Hexamethylenetetramine Hydrosulphite (hydrosulfite) Sodium Hydrosulphite
Ichthyol

ĸ

Kauri Gum	 Copal, Gum	
Kieselguhr	 Tripoli Diatomaceous	Earth

L

Lanum	Lanolin
Lead Chromate	Chrome Yellow
Lead Sulfate, Basic	Whitelead, Sublimed
Lemon, Salts of	Potassium Binoxalate
Licorice	Glycyrrhiza
Ligroin, Light	Petroleum Ether
Lime	Calcium Oxide
Lime, Slaked	Calcium Hydroxide
Limestone	Calcium Carbonate
Litharge	Lead Monoxide
Liver of Sulphur	Potassium Sulphide
Lunar Caustic	
Lye	Sodium Hydroxide

M

Magnesium, Calcined	Magnesium Oxide
Magnesium Silicate	Talcum
Maize Oil	Corn Oil
Malt Sugar	Maltose
Metol	Methyl-para-aminophenol Sulphate
Microcosmic Salt	Sodium Ammonium Phosphate

Milk Sugar Mineral Pitch Minium Mirbane Oil Muriatic Acid Myrtle Wax	Asphalt Lead Oxide (red) Nitrobenzol Hydrochloric Acid
Myrtle Wax	Bayberry Wax

N

Naphtha, Solvent	Coal Tar Naphtha
Naples Yellow	
Nickel Salts, Double	
Nickel Salts, Single	Nickel Sulphate
Niter	
Niter Cake	Sodium Bisulphate
Nitrocellulose (soluble cotton)	Pyroxylin

0

Oleic Acid	Red Oil
Olein	. Glyceryl Tri-oleate (natural)
Oleum	Sulphuric Acid (fuming)
Olive Oil	Sweet Oil
Orange Mineral	Orange Red Lead Oxide
Orpiment	Arsenous Sulphide (yellow)

P

Paraffin Oil Paris White	Mineral Oil
Paramii On	Petrolatum, Liquid
Paris White	Whiting
Pearl Ash	.Potassium Carbonate
Petrol	
Petrolatum	
Plaster of Paris	
Potassium Bicarbonate	
Prussian Blue	.Ferric Ferrocyanide
Prussiate of Potash, Red	
Prussiate of Potash, Yellow	.Potassium Ferrocyanide
Prussic Acid	.Hydrocyanic Acid
Pyramidon	.Amidopyrine
Pyrethrum	.Insect Flowers (powdered)
Pyroligneous Acid	·Wood Vinegar

Q

Quicklime																	Cal	ciur	n	Oz	cide	Э
Quicksilver	•			•	•		•		•		•	•	•		•	•	Mei	ccur	у			

R

Red Oxide	Ferric	Oxi	de, Red	
Rochelle Salt	Potass	ium	Sódium	Tartrate
Rottenstone	Tripol	i		

5

Saccharine	Glucoside
Sal Ammoniac	Ammonium Chloride
Sal Soda	Sodium Carbonate, Hydrated
Salad Oil	Cottonseed Oil
Salt	Sodium Chloride
Salt Cake	

Saltpeter Potassium Nitrate Scale Wax Paraffin Wax (low melting) Silica Silicon Dioxide Sod Oil Degras Soda Ash Sodium Carbonate, Anhydrous Sodium Bisulphite Sodium Acid Sulphate Sodium Phosphate, Dibasic Disodium Phosphate Sodium Phosphate, Monobasic Monosodium Phosphate Sodium Phosphate, Tribasic Trisodium Phosphate Sodium Phosphate Hypp Sperm Oil Whale Oil Spirits of Turpentine Turpentine Stannous Chloride Tin Crystals Stearin Tristearin Storax Styrax Sucrose Cane Sugar Sugar of Lead Lead Acetate Sulfonated Castor Oil Olive Oil Foots Sulphuric Acid Oil of Vitriol Sulphuric Ether Ether
т
TNT Trinitrotoluene Tartar Emetic Antimony Potassium Tartrate Tetralin Tetrahydro Naphthalene Theobroma Oil Cacao Butter Titanium Dioxide Titanium Oxide Toluene Toluel Triacetin Glycerol Triacetate Trinitrophenol Picric Acid
V
Verdigris
w
Whale Oil Train Oil White Arsenic Arsenic Trioxide White Bole Kaolin White Lead Lead Carbonate, Basic White Metal Babbitt Metal White Wax Beeswax (bleached) Whiting Chalk, Refined Wintergreen Oil, Synthetic Methyl Salicylate Wood Alcohol Methyl Alcohol
Y
Yacca GumAccroides Gum
z
Zinc White

TRADE NAMED CHEMICALS

During the past few years, the practice of marketing raw materials, under names which in themselves are not descriptive chemically of the products they represent, has become very prevalent. No modern book of formulae could justify its claims either to completeness or modernity without numerous formulae containing these so-called "Trade Names."

Without wishing to enter into any discussion regarding the justification of "Trade Names," the Editors recognize the tremendous service rendered to commercial chemistry by manufacturers of "Trade Name" products, both in the physical data supplied and the formulation suggested.

Deprived of the protection afforded their products by this system of nomenclature, these manufacturers would have been forced to stand helplessly by while the fruits of their labor were being filched from them by competitors who, unhampered by expenses of research, experimentation and promotion, would be able to produce something "just as good" at prices far below those of the original producers.

That these competitive products were "just as good" solely in the minds of the imitators would only be evidenced in costly experimental work on the part of the purchaser and, in the meantime irreparable damage would have been done, to the truly ethical product. It is obvious, of course, that under these circumstances, there would be no incentive for manufacturers to develop new materials.

Because of this, and also because the "Chemical Formulary" is primarily concerned with the physical results of compounding rather than with the chemistry involved, the Editors felt that the inclusion of formulae containing various trade name products would be of definite value to the producer of finished chemical materials. If they had been left out many ideas and processes would have been automatically eliminated.

As a further service a list of the better known "trade name" products is appended together with the suppliers of these materials. The number after each trade name refers to the supplier given below with the corresponding number.

TRADE NAMES

	A consists 156
A	Ascarite
A-Syrup	Astruian 6 Atrapol
Abalyn 79	Aurosal
Abopon	Aurosai
Accelerator 808	Avonac
Accelerator 833	В
Acetoin	Badex151
Acidolene	Bakelite
Acto	Bardol
Adheso Wax	Barretan
A.D.M. No. 100 Oil 10	Beckacite
Aerogel101	Beckolin
Agerite Powder163	Beckosol
Akcocene 6	Bensapol
Alba-Floc	Beutene
Albasol106	Blandol
Albatex 38	Blendene
Albertol	Bludtan 33
Albinol136	Bordow 49
Albolith110	Borol 50
Albone "C" 51	Bromo "Acid"
Albusol	Brosco
Aldehol 87	Butalyde 42
Aldol181	Butyl Carbitol
Alkanol	Butyl Cellosolve 28
Alloxan 20	С
Aloxite	
Alphasol	Cadalyte 73
Altax163	Cadmolith
Alugel104	Calcoloid
Alugel 104 Amberette 154	Calcoloid 25 Calcene 41
Alugel 104 Amberette 154 Amberol 125	Calcoloid25Calcene41Calgon22
Alugel 104 Amberette 154 Amberol 125 Ambreno 51	Calcoloid 25 Calcene 41 Calgon 22 Calorite 148
Alugel 104 Amberette 154 Amberol 125 Ambreno 51 Amco Acetate 88	Calcoloid 25 Calcene 41 Calgon 22 Calorite 148 Captax 165
Alugel 104 Amberette 154 Amberol 125 Ambreno 51 Amco Acetate 88 Amandol 51	Calcoloid 25 Calcene 41 Calgon 22 Calorite 148 Captax 165 Carbitol 28
Alugel 104 Amberette 154 Amberol 125 Ambreno 51 Amco Acetate 88 Amandol 51 Amidine 26	Calcoloid 25 Calcene 41 Calgon 22 Calorite 148 Captax 165 Carbitol 28 Carboxide 28
Alugel 104 Amberette 154 Amberol 125 Ambreno 51 Amco Acetate 88 Amandol 51 Amidine 26 Anchoracel 2p 7	Calcoloid 25 Calcene 41 Calgon 22 Calorite 148 Captax 165 Carbitol 28 Carboxide 28 Caseo 30
Alugel 104 Amberette 154 Amberol 125 Ambreno 51 Amco Acetate 88 Amandol 51 Amidine 26 Anchoracel 2p 7 Anhydrone 14	Calcoloid 25 Calcene 41 Calgon 22 Calorite 148 Captax 165 Carbitol 28 Carboxide 28 Casco 30 Catalpo 102
Alugel 104 Amberette 154 Amberol 125 Ambreno 51 Amco Acetate 88 Amandol 51 Amidine 26 Anchoracel 2p 7 Anhydrone 14 Ansol 161	Calcoloid 25 Calcene 41 Calgon 22 Calorite 148 Captax 165 Carbitol 28 Carboxide 28 Casco 30 Catalpo 102 CCH 98
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Alugel 104 Amberette 154 Ambreno 125 Amoreno 51 Amco Acetate 88 Amandol 51 Amidine 26 Anchoracel 2p 7 Anhydrone 14 Ansol 161 Antidolorin 58 Apcothinner 8	Calcoloid 25 Calcene 41 Calgon 22 Calorite 148 Captax 165 Carbitol 28 Carboxide 28 Casco 30 Catalpo 102 CCH 98 Celascour 3 Celite 85
Alugel 104 Amberette 154 Ambreno 125 Amco Acetate 88 Amandol 51 Amidine 26 Anchoracel 2p 7 Anhydrone 14 Ansol 161 Antidolorin 58 Apcothinner 8	Calcoloid 25 Calcene 41 Calgon 22 Calorite 148 Captax 165 Carbitol 28 Carboxide 28 Casco 30 Catalpo 102 CCH 98 Celascour 3 Celite 85 Cellosolve 28
Alugel 104 Amberette 154 Amberol 125 Ambreno 51 Amco Acetate 88 Amandol 51 Amidine 26 Anchoracel 2p 7 Anhydrone 14 Ansol 161 Antidolorin 58 Apcothinner 8 Aqualoid 86	Calcoloid 25 Calcene 41 Calgon 22 Calorite 148 Captax 165 Carbitol 28 Carboxide 28 Casco 30 Catalpo 102 CCH 98 Celascour 3 Celite 85 Cellosolve 28 Censteric 32
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Alugel 104 Amberette 154 Amberol 125 Ambreno 51 Amco Acetate 88 Amandol 51 Amidine 26 Anchoracel 2p 7 Anhydrone 14 Ansol 161 Antidolorin 58 Apcothinner 8 Aqualoid 86 Aquamel 70 Aquapel 114	Calcoloid 25 Calcene 41 Calgon 22 Calorite 148 Captax 165 Carbitol 28 Carboxide 28 Casco 30 Catalpo 102 CCH 98 Celascour 3 Celite 85 Cellosolve 28 Censteric 32 Cerelose 44 Cereps 170
Alugel 104 Amberette 154 Amberol 125 Ambreno 51 Amco Acetate 88 Amandol 51 Amidine 26 Anchoracel 2p 7 Anhydrone 14 Ansol 161 Antidolorin 58 Apcothinner 8 Aqualoid 86 Aquamel 70 Aquapel 114 Aquarome 55	Calcoloid 25 Calcene 41 Calgon 22 Calorite 148 Captax 165 Carbitol 28 Carboxide 28 Casco 30 Catalpo 102 CCH 98 Celascour 3 Celite 85 Cellosolve 28 Censteric 32 Cerelose 44 Cereps 170 Ceresalt 53
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Alugel 104 Amberette 154 Amberol 125 Ambreno 51 Amco Acetate 88 Amandol 51 Amidine 26 Anchoracel 2p 7 Anhydrone 14 Ansol 161 Antidolorin 58 Apcothinner 8 Aqualoid 86 Aquapel 114 Aquarome 55 Aquasol 6 Arapali 129 Araskleen 101 Archer-Daniels No. 635 10	Calcoloid 25 Calcene 41 Calgon 22 Calorite 148 Captax 165 Carbitol 28 Carboxide 28 Casco 30 Catalpo 102 CCH 98 Celascour 3 Celite 85 Cellosolve 28 Censteric 32 Cerelose 44 Cereps 170 Ceresalt 53 Chlorex 28
Alugel 104 Amberette 154 Amberol 125 Ambreno 51 Amco Acetate 88 Amandol 51 Amidine 26 Anchoracel 2p 7 Anhydrone 14 Ansol 161 Antidolorin 58 Apcothinner 8 Aqualoid 86 Aquamel 70 Aquapel 114 Aquarome 55 Aquasol 6 Arapali 129 Araskleen 101 Archer-Daniels No. 635 10 Archer-Daniels-Midland Oil 10	Calcoloid 25 Calcene 41 Calgon 22 Calorite 148 Captax 165 Carbitol 28 Carboxide 28 Casco 30 Catalpo 102 CCH 98 Celascour 3 Celite 85 Cellosolve 28 Cersteric 32 Cerelose 44 Cereps 170 Ceresalt 53 Chlorex 28 Chlorasol 28 Chremnitz White 56 Cinchophen 25
Alugel 104 Amberette 154 Amberol 125 Ambreno 51 Amco Acetate 88 Amandol 51 Amidine 26 Anchoracel 2p 7 Anhydrone 14 Ansol 161 Antidolorin 58 Apcothinner 8 Aqualoid 86 Aquanel 70 Aquapel 114 Aquasol 6 Arapali 129 Araskleen 101 Archer-Daniels No. 635 10 Archer-Daniels-Midland Oil 10 Aridex 51	Calcoloid 25 Calcene 41 Calgon 22 Calorite 148 Captax 165 Carbitol 28 Carboxide 28 Casco 30 Catalpo 102 CH 98 Celascour 3 Celite 85 Cellosolve 28 Censteric 32 Cerelose 44 Cereps 170 Ceresalt 53 Chlorex 28 Chlorasol 28 Chremnitz White 56 Cinchophen 25 Coblac 19
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Alugel 104 Amberette 154 Amberol 125 Ambreno 51 Amco Acetate 88 Amandol 51 Amidine 26 Anchoracel 2p 7 Anhydrone 14 Ansol 161 Antidolorin 58 Apcothinner 8 Aqualoid 86 Aquapel 114 Aquarome 55 Aquasol 6 Arapali 129 Araskleen 101 Archer-Daniels No. 635 10 Archer-Daniels-Midland Oil 10 Aridex 51 Arcochlor 153 Arosol 64	Calcoloid 25 Caleene 41 Calgon 22 Calorite 148 Captax 165 Carbitol 28 Carboxide 28 Casco 30 Catalpo 102 CCH 98 Celascour 3 Celite 85 Cellosolve 28 Censteric 32 Cerelose 44 Cereps 170 Ceresalt 53 Chlorex 28 Chlorasol 28 Chremnitz White 56 Cinchophen 25 Coblac 19 Cominol 43 Coppercide 83
Alugel 104 Amberette 154 Amberol 125 Ambreno 51 Amco Acetate 88 Amandol 51 Amidine 26 Anchoracel 2p 7 Anhydrone 14 Ansol 161 Antidolorin 58 Apcothinner 8 Aqualoid 86 Aquapel 114 Aquarome 55 Aquasol 6 Arapali 129 Araskleen 101 Archer-Daniels No. 635 10 Archer-Daniels-Midland Oil 10 Aridex 51 Arosol 64 Artisil 134	Calcoloid 25 Calcene 41 Calgon 22 Calorite 148 Captax 165 Carbitol 28 Carboxide 28 Casco 30 Catalpo 102 CCH 98 Celascour 3 Celite 85 Cellosolve 28 Censteric 32 Cerelose 44 Cereps 170 Ceresalt 53 Chlorex 28 Chlorex 28 Chremnitz White 56 Cinchophen 25 Coblac 19 Cominol 43 Coppercide 83 Cosmic Black 158
Alugel 104 Amberette 154 Amberol 125 Ambreno 51 Amco Acetate 88 Amandol 51 Amidine 26 Anchoracel 2p 7 Anhydrone 14 Ansol 161 Antidolorin 58 Apcothinner 8 Aqualoid 86 Aquapel 114 Aquarome 55 Aquasol 6 Arapali 129 Araskleen 101 Archer-Daniels No. 635 10 Archer-Daniels-Midland Oil 10 Aridex 51 Arcochlor 153 Arosol 64	Calcoloid 25 Caleene 41 Calgon 22 Calorite 148 Captax 165 Carbitol 28 Carboxide 28 Casco 30 Catalpo 102 CCH 98 Celascour 3 Celite 85 Cellosolve 28 Censteric 32 Cerelose 44 Cereps 170 Ceresalt 53 Chlorex 28 Chlorasol 28 Chremnitz White 56 Cinchophen 25 Coblac 19 Cominol 43 Coppercide 83

Cromodine	1
Cryptone	•
Cumar	Idalol
Cyclamal	Igepon
Cycline	IG Wax O
Cymanol	Indian Red
Cymanor	Indigisols
D	Indur124
*	Isolene
Darco 45	
Diamond K Linseed Oil145	J
Dionin	T
Discolite	Jasmogene
Disperso	К
Distoline	K.
Duolith 90	Kalite163
Duphax146	Karo
Duphonol	
DuPont Rubber Red 51	Kellogg Kuo .145 Kellogg Varnish Oil .145
Durez 68	Kerol
	Kilfoam 4
E	Kolineum 89
	Kopol
Eastman Products 52	Koreon103
Elaine 54	Kryocide
Erio Chrome Dyes	ixiyocide
Esterol	. L
Estersol	
Ethox	Lactol Spirits
Ethyl Parasept179	Lacquer Blue 9
Ethyl Protol 48	Lanette Wax
Eulan 65	Laurex108
	Le Page's Cement
F	Leukonin 77
	Leukonin 77 Lewisol 92
Factolac 81	Leukonin 77
Factolac	Leukonin 77 Lewisol 92 Lindol 31 Lohrinol 70
Factolac 81 Falba Absorption Base 119 Feectol 131	Leukonin 77 Lewisol 92 Lindol 31 Lohrinol 70 Lucidol 94
Factolac 81 Falba Absorption Base 119 Feectol 131 Fer-ox 173	Leukonin 77 Lewisol 92 Lindol 31 Lohrinol 70
Factolac 81 Falba Absorption Base 119 Feectol 131 Fer-ox 173 Ferrox 150	Leukonin 77 Lewisol 92 Lindol 31 Lohrinol 70 Lucidol 94 Lysol 91
Factolac 81 Falba Absorption Base 119 Feectol 131 Fer-ox 173 Ferrox 150 Fixalt 101	Leukonin 77 Lewisol 92 Lindol 31 Lohrinol 70 Lucidol 94
Factolac 81 Falba Absorption Base 119 Feectol 131 Fer-ox 173 Ferrox 150 Fixalt 101 Flexoresin 70	Leukonin 77 Lewisol 92 Lindol 31 Lohrinol 70 Lucidol 94 Lysol 91
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Factolac 81 Falba Absorption Base 119 Feectol 131 Fer-ox 173 Ferrox 150 Fixalt 101 Flexoresin 70	Leukonin
Factolac 81 Falba Absorption Base 119 Feectol 131 Fer-ox 173 Ferrox 150 Fixalt 101 Flexoresin 70 Fyrex 166	Leukonin
Factolac 81 Falba Absorption Base 119 Feectol 131 Ferox 173 Ferrox 150 Fixalt 101 Flexoresin 70 Fyrex 166 G Gardinol 51	Leukonin 77 Lewisol 92 Lindol 31 Lohrinol 70 Lucidol 94 Lysol 91 M Mapico 19 Mellittis 69 Merpentine 51 Methyl Cellosolve 28 Metso 120
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Factolac 81 Falba Absorption Base 119 Feectol 131 Fer-ox 173 Ferrox 150 Fixalt 101 Flexoresin 70 Fyrex 166 G Gardinol Gastex 62 Gelya 140	Leukonin 77 Lewisol 92 Lindol 31 Lohrinol 70 Lucidol 94 Lysol 91 M Mapico 19 Mellittis 69 Merpentine 51 Methyl Cellosolve 28 Metso 120
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Factolac 81 Falba Absorption Base 119 Feectol 131 Ferox 173 Ferrox 150 Fixalt 101 Flexoresin 70 Fyrex 166 G Gardinol 51 Gastex 62 Gelva 140 Gilsonite 15 Glycopon 70 Glytosterin 70 Glytptal 66 Guai-a-phene 40 Guantal 131 H Halowax Hercusol 79 Hydralite C 65	Leukonin
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SUPPLIERS OF TRADE NAME CHEMICALS

- Acheson Graphite Corp., Niagara Falls, N. Y.
- Acheson Graphite Corp., Niagara Falls, N. Y.
 Advance Solvents & Chem. Corp., New York City
 American Aniline Products, Inc., New York City
 American Chem. Prod. Co., Rochester, N. Y.
 American Colloid Co., Chicago, Ill.
 American Cyanamid & Chem. Co., New York City
 Anchor Chem. Co., Manchester, England
 Anderson Prichard Oil Corp., Oklahoma City, Okla.
 Ansbacher-Siegle Corp., Rosebank, N. Y.
 Archer-Daniels-Midland Co., Minneapolis, Minn.
 Arkansas Co.. New York City

- 11. Arkansas Co., New York City

- Arkanias So., New York City
 Atlantic Refining Co., Phila., Pa.
 Bakelite Corp., New York City
 Baker, J. T. Chem. Co., Phillipsburg, N. J.
 Barber Asphalt Co., Phila., Pa.
 Barrett Co., New York City

- 17. Beck, Koller & Co., Detroit, Mich.

- 18. Bick & Co., Inc., Reading, Pa.

 19. Binney & Smith, New York City

 20. British Drug Houses, Ltd., London, England

 21. Bud Aromatic Chem. Co., Inc., New York City

- 21. Bud Aromatic Chem. Co., Inc., New York City
 22. Buromin Corp., Pittsburgh, Pa.
 23. Bush, W. J. & Co. Inc., New York City
 24. Cabot, Godfrey L. Inc., Boston, Mass
 25. Calco Chem. Co., Bound Brook, N. J.
 26. Campbell, John & Co., New York City
 27. Carbic Color & Chem. Co., New York City
 28. Carbide & Carbon Chem. Corp., New York City
 29. Carborundum Co., Niagara Falls, N. Y.
 30. Casein Mfg. Co., New York City
 31. Celluloid Corp., Newark, N. J.
 32. Century Stearic Acid & Candle Wks., New York City
 33. Champion Fibre Co., Canton, No. Car.

- 32. Century Stearic Acid & Candle Wks., New York Cit
 33. Champion Fibre Co., Canton, No. Car.
 34. Chaplin-Bibbo, New York City
 35. Chemical & Pigment Co., Inc., Scranton, Pa.
 36. Chemical Solvents Inc., New York City
 37. Chesebrough Mfg. Co., New York City
 38. Ciba Co., Inc., New York City
 39. Colgate-Palmolive-Peet Co., Jersey City, N. J.
 40. Colledge, E. W., Inc., Cleveland, O.
 41. Columbia Alkali Corp., New York City
 42. Commercial Solvents Corp., Terre Haute, Ind.
 43. Commonwealth Color & Chem. Co., Brooklyn, N. Y.
 44. Corn Products Refining Co., New York City
 45. Darco Sales Corp., New York City
 46. Deep Rock Oil Corp., Chicago, Ill.

- 46. Deep Rock Oil Corp., Chicago, Ill. 47. Dennis, Martin & Co., Newark, N. J.

- 47. Dennis, Martin & Co., Newark, N. J.

 48. Dodge & Olcott Co., New York City

 49. Dow Chem. Co., Midland, Mich.

 50. Ducas, B. P. Co., New York City

 51. DuPont, E. I., de Nemours & Co., Wilmington, Del.

 52. Eastman Kodak Co., Rochester, N. Y.

 53. Economic Materials Co., Chicago, Ill.

 54. Emery Industries, Inc., Cincinnati, O.

 55. Felton Chem. Co., Brooklyn, N. Y.

 56. Fezandié and Sperrlé. Inc., New York City

- 56. Fezandié and Sperrlé, Inc., New York City
- 57. Fougera, E. & Co., New York City
- 58. Franco-Amer. Chem. Works, Carlstadt, N. J.

59. Fries Bros., New York City

60. Fritzchie Bros., New York City

61. Geigy Co. Inc., New York City

62. General Atlas Carbon Co., New York City 63. General Chemical Co., New York City

64. General Drug Co., New York City65. General Dyestuffs Corp., New York City

66. General Electric Co., Schenectady, N. Y.

67. General Naval Stores Co., New York City

67. General Navai Stores Co., New York City
68. General Plastics Inc., No. Tonawanda, N. Y.
69. Givaudan-Delawanna, Inc., New York City
70. Glyco Products Co., Inc., New York City
71. Goldschmidt Corp., New York City
72. Goodyear Tire & Rubber Co., Akron, O.
73. Grasselli Chem. Co., Cleveland, O.
74. Greef, R. W. & Co., Inc., New York City
75. Hall, C. P. & Co., Akron, O.
76. Haloway Corp. New York City

76. Halowax Corp., New York City 77. Harshaw Chem. Co., Cleveland, O.

78. Heine & Co., New York City 79. Hercules Powder Co., Wilmington, Del.

80. Hooker Electro-Chem. Co., New York City

81. Hopkins, J. L. & Co., New York City 82. Industrial Chem. Sales Co., New York City 83. Innis, Speiden & Co., New York City

84. International Pulp Corp., New York City 85. Johns-Manville Corp., New York City

86. Jungmann & Co., New York City

87. Kay-Fries Chemicals, Inc., New York City 88. Kessler Chem. Corp., New York City 89. Koppers Products Co., Pittsburgh, Pa.

90. Krebs Pigment & Color Corp., Newark, N. J. 91. Lehn & Fink Corp., New York City

92. Lewis, John D., Inc., Providence, R. I. 93. Liquid Carbonic Corp., Chicago, Ill. 94. Lucidol Corp., Buffalo, N. Y.

95. Magnus, Mabee & Reynard, Inc., New York City 96. Mallinckrodt Chem. Works, St. Louis, Mo. 97. Martin, Dennis Co., Newark, N. J. 98. Mathieson Alkali Co., New York City 99. McCormick & Co., Baltimore, Md.

100. Merck & Co. Inc., New York City 101. Monsanto Chem. Works, St. Louis, Mo.

102. Moore-Munger, New York City

103. Mutual Chem. Co. of Amer., Newark, N. J. 104. National Aluminate Corp., Chicago, Ill. 105. National Aniline & Chem. Co., Buffalo, N. Y.

106. National Oil Products Co., Harrison, N. J. 107. National Rosin Oil & Size Co., New York City

108. Naugatuck Chem. Co., New York City

109. Neville Co., Pittsburgh, Pa. 110. New Jersey Zinc Sales Co., New York City

111. Nulomoline Co., New York City 112. Nuodex Products, Inc., Newark, N. J. 113. Onyx Oil & Chem. Co., Passaic, N. J.

113. Onyx Oll & Chem. Co., Passaic, N. J.
114. Papermakers' Chem. Corp., Wilmington, Del.
115. Paramet Chem. Corp., Long Island City, N. Y.
116. Penick, S. B. & Co., New York City
117. Penn. Alcohol Corp., Phila., Pa.
118. Penn. Salt Mfg. Co., Phila., Pa.
119. Pfaltz & Bauer, Inc., New York City
120. Phila. Quartz Co., Phila., Pa.
121. Plymouth Organic Labs., New York City
122. Pylam Products Co., New York City
123. Rauh, Robert Inc., Newark, N. J.

123. Rauh, Robert Inc., Newark, N. J.

124. Reilly Tar & Chem. Corp., Indianapolis, Ind. 125. Resinous Prod. & Chem. Co., Philadelphia, Pa.

126. Resinox Corp., New York City

- 127. Revertex Corp., New York City
- 128. Robeson Process Co., New York City 129. Rohm-Hass Chem. Co., Philadelphia, Pa. 130. Royce Chem. Co., Carlton Hill, N. J.
- 131. Rubber Service Labs. Co., Akron, O.
- 132. Russia Cement Co., Gloucester, Mass. 133. Salomon, L. A. & Bro., New York City 134. Sandoz Chem. Works, New York City
- 135. Scholler Bros., Inc., Philadelphia, Pa.
- 136. Schliemann Co., Inc., New York City 137. Scott, Bader & Co., London, England 138. Seeley & Co., New York City
- 139. Sharples Solvents Corp., Philadelphia, Pa. 140. Shawinigan, Ltd., New York City
- 141. Sherwood Petroleum Co., Brooklyn, N. Y.
- 141. Silver, Geo., Import Co., New York City 142. Silver, Geo., Import Co., New York City 143. Sonneborn, L. Sons, New York City 144. Southwark Mfg. Co., Camden, N. J. 145. Spencer-Kellogg Co., New York City
- 146. Stamford Rubber Supply Co., Stamford, Conn.
- 147. Stanco, Inc., New York City 148. Standard Oil Co. of Calif., San Francisco, Cal. 149. Standard Oil Co. of New Jersey, New York City
- 150. Stauffer Chem. Co., New York City
- 151. Stein-Hall & Co., Inc., New York City 152. Sun Oil Co., Philadelphia, Pa.
- 153. Swann Chem. Corp., Birmingham, Ala.
- 154. Synfleur Scientific Labs., Monticello, N. Y. 155. Texas Mining & Smelting Co., Laredo, Texas
- 156. Thomas, Arthur H., Co., Philadelphia, Pa.
 157. Titanium Pigments Co., New York City
 158. Uhlich, Paul Co., New York City
- 159. United Color & Pigment Co., Inc., Newark, N. J. 160. United States Gypsum Co., Chicago, Ill.
- 161. United States Industrial Chem. Co., Inc., New York City
- 162. Van-Ameringen Haebler, Inc., New York City
- 163. Vanderbilt, R. T. Co., Inc., New York City 164. Varcum Chem. Corp., Niagara Falls, N. Y. 165. Verley, Albert & Co., Chicago, Ill.
- 166. Victor Chem. Works, Chicago, Ill.
- 167. Virginia Smelting Co., W. Norfolk, Va. 168. Vultex Corp. of America, Cambridge, Mass.
- 169. Wallerstein Co., Inc., New York City 170. Welch, Holme & Clark Co., Inc., New York City
- 171. Whittaker, Clark & Daniels, Inc., New York City 172. Will & Baumer Candle Co., New York City
- 173. Wishnick-Tumpeer, Inc., New York City 174. Woburn Degreasing Co. of N. J., Harrison, N. J.
- 175. Wolf, Jacques & Co., Passaic, N. J.
- 176. Amer. Chemical Paint Co., Rochester, N. Y.
- 177. Baker & Co., Inc., Newark, N. J. 178. Chemical & Pigment Co., Baltimore, Md.
- 179. Heyden Chem. Works, New York, N. Y. 180. Kali Mfg. Co., Philadelphia, Pa.
- 181. Niacet Chem. Corp., Niagara Falls, N. Y. 182. Proctor & Gamble, Cincinnati, Ohio.
- 183. Pure Calcium Products Co., Gainesville, O. 184. Van Schaack Bros. Chem. Co., Chicago, Ill.
- 185. Wyodak Chem. Co., Cleveland, O.

WHERE TO BUY CHEMICALS

Abietic Acid

Hercules Powder Co., New York, N. Y.

Accelerators, Vulcanization

Rubber Service Labs., Inc., Akron, O.

Acetamide

Amer. Chemical Products Co., Rochester, N. Y.

Acetic Acid

The Cleveland-Cliffs Iron Co., Cleveland, Ohio

Acetic Anhydride

American-British Chemical Supplies, Inc., New York, N. Y.

Acetone

W. S. Gray Co., New York, N. Y.

Acetphenetidin

Merck & Co., Inc., Rahway, N. J.

Acetyl Salicylic Acid

Monsanto Chemical Co., St. Louis, Mo.

Acids, Fatty

Arthur C. Trask Co., Chicago, Ill.

Acriflavine

Abbott Laboratories, North Chicago, Ill.

A aar

American Agar Co., Inc., San Diego, Calif.

Albumen Stein, Hall & Co., Inc., New York, N. Y.

Alcohol, Denatured

Rogers & McClellan, Boston, Mass. L. R. Van Allen & Co., Chicago, Ill.

Alcohol, Pure

U. S. Industrial Alcohol Co., New York, N. Y.

Allealies

Columbia Alkali Corp., New York, N. Y.

Alkaloids

Merck & Co., Inc., Rahway, N. J.

Alkanet

J. L. Hopkins & Co., New York, N. Y.

Almond Oil

Magnus, Mabee & Reynard, Inc., New York, N.

Aloes

Peck & Velsor, New York, N. Y.

Alpha Naphthol

Hord Color Products, Sandusky, O.

Alumina

Aluminum Co. of America, Pittsburgh, Pa.

Aluminum

Aluminum Co. of America, Pittsburgh, Pa.

Aluminum Hydrate
Ceramic Color & Chem. Mfg. Co., New Brighton, Pa.

Alums
The Grasselli Chemical Co., Cleveland, O.

Aluminum Acetate
Niacet Chemicals Corp., Niagara Falls, N. Y.

Aluminum Bronze Powder
U. S. Bronze Powder Works, Inc., New York, N. Y.

Aluminum Chloride (Solution, Crystals and Anhydrous)
The Calco Chemical Co., Bound Brook, N. J.

Aluminum Stearate
Franks Chemical Products Co., Inc., Brooklyn, N. Y.

Aminostearin
Glyco Products Co., Inc., New York, N. Y.

Ammonia
Nat'l Ammonia Co., Inc., Philadelphia, Pa.

Ammonium Bifluoride
The Harshaw Chemical Co., Cleveland, C.

Ammonium Carbonate
Wishnick-Tumpeer, Inc., New York, N. Y.

Ammonium Chloride Pennsylvania Salt Mfg. Co., Inc., Philadelphia, Pa.

Ammonium Linoleate
Glyco Products Co., Inc., New York, N. Y.

Ammonium Nitrate
Garrigues, Stewart & Davies, Inc., New York, N. Y.

Ammonium Oleate
Glyco Products Co., Inc., New York, N. Y.

Ammonium Persulphate
Buffalo Electro Chemical Co., Inc., Buffalo, N. Y.

Ammonium Phosphate Swann Chemical Co., New York, N. Y.

Ammonium Sulphate H. J. Baker & Bro., New York, N. Y.

Ammonium Stearate
Glyco Products Co., Inc., New York, N. Y.

Amyl Acetate
Chemical Solvents, Inc., New York, N. Y.

Aniline Dyes
Experimenter's Supply Co., New York, N. Y.

Aniline Oil
Dow Chemical Co., Midland, Michigan

Antimony
C. Tennant & Sons Co. of N. Y., New York, N. Y.

Antimony Chloride
Seldner & Enequist, Inc., Brooklyn, N. Y.

Antimony Oxide
O. Hommel Co., Pittsburgh, Pa.

Antimony Sulphide
Foote Mineral Co., Philadelphia, Pa.

Anti Oxidants
Givaudan-Delawanna, Inc., New York, N. Y.

Arsenic

Amer. Smelting & Refining Co., New York, N. Y.

Asbestos

Powhatan Mining Corp., Woodlawn, Baltimore, Md.

Asphalt

The Barber Asphalt Co., Philadelphia, Pa.

Asphaltum

Allied Asphalt & Mineral Corp., New York, N. Y.

Balsams

James B. Horner, Inc., New York, N. Y.

Barium Carbonate

Barium Reduction Corp., Charleston, W. Va.

Barium Nitrate

C. W. Campbell Co., Inc., New York, N. Y.

Barium Peroxide

Barium Reduction Corp., Charleston, W. Va. Barium Sulphate

C. P. De Lore Co., St. Louis, Mo.

Barium Sulphide
Chicago Copper & Chemical Co., Blue Island, Ill.

Barutes

Bradley & Baker, New York, N. Y. Nat'l Pigments & Chemical Co., St. Louis, Mo.

Basic Colors

Amer. Aniline Products, Inc., New York, N. Y.

Bayberry Wax

The W. H. Bowdlear Co., Syracuse, N. Y.

Beeswax

A. C. Drury & Co., Inc., Chicago, Ill. Theodor Leonhard Wax Co., Inc., Haledon, Paterson. N. J.

Bentonite

Amer. Colloid Co., Chicago, Ill. Silica Products Co., Kansas City, Mo. The Wyodak Chemical Co., Cleveland, Ohio

Benzaldehyde

Heyden Chem. Corp., New York, N. Y.

Benzidine

General Aniline Works, Inc., New York, N. Y.

Benzine

Amer. Mineral Spirits Co., New York, N. Y.

Benzocaine

Abbott Laboratories, No. Chicago, Ill.

Benzoic Acid

Carus Chemical Co., Inc., La Salle, Ill.

Benzol

The Barrett Co., New York, N. Y.

Benzoyl Peroxide

Lucidol Corp., Buffalo, N. Y.

Benzyl Cellulose

Advance Solvents & Chem. Corp., New York, N. Y.

Bergamot Oil

Orbis Products Corp., New York, N. Y.

Beryllium

Belmont Smelting & Refining Wks., Inc., Brooklyn, N. Y.

Beryllium and Its Salts

Beryllium Corp. of America, New York, N. Y.

Beta Naphthol

The Calco Chemical Co., Bound Brook, N. J.

Bismuth

Cerro de Pasco Copper Corp., New York, N. Y.

Bismuth Subnitrate

The New York Quinine & Chemical Wks., Inc., Brooklyn, N. Y.

Blanc Fixe

Adolph Hurst & Co., Inc., New York, N. Y.

Bleaching Powder

Electro Bleaching Gas Co., New York, N. Y.

Blood Albumen

Morningstar, Nicol, Inc., New York, N. Y.

Bone Ash

Denver Fire Clay Co., Denver, Colorado

Bone Black

Siemon Colors, Inc., Newark, N. J.

Bone Glue

Darling & Co., Chicago, Ill.

Bone Oil

Texas Chemical Co., Houston, Texas

Borax

American Potash & Chem. Corp., New York, N. Y.

Bordeaux Mixture

Mechling Bros. Chem. Co., Camden, N. J.

Boric Acid

Borax Union, Inc., San Francisco, Calif.

Botanical Products

S. B. Penick & Co., New York, N. Y.

Bromine

J. Q. Dickinson & Co., Malden, W. Va.

Bromo-Fluorescein

Glyco Products Co., Inc., New York, N. Y.

Bronze Powder

B. K. Drakenfeld & Co., New York, N. Y.

Burgundy Pitch

Geo. H. Lincks, New York, N. Y.

Butyl Acetate

Commercial Solvents Corp., New York, N. Y. Publicker, Inc., Philadelphia, Pa.

Butyl Aldehyde

Commercial Solvents Corp., Terre Haute, Ind.

Butyl Alcohol (Normal)

Publicker, Inc., Philadelphia, Pa.

Butyl Propionate
C. P. Chemical Solvents, Inc., New York, N. Y.

Butyric Ether

The Northwestern Chemical Co., Wauwatosa, Wisconsin

Butyl Stearate

Kessler Chem. Corp., New York, N. Y.

Cadmium

U. S. Smelting, Refining & Mining Co., New York, N. Y.

Cajuput Oil

D. W. Hutchinson & Co., New York, N. Y.

Calcium Arsenate

Bowker Chemical Corp., New York, N. Y. Chipman Chemical Co., Inc., Bound Brook, N. J.

Calcium Carbonate

Limestone Products Corp. of Amer., Newton, N. J.

Calcium Carbonate (Precipitated) Merck & Co., Inc., Rahway, N. J.

Calcium Chloride

Michigan Alkali Co., New York, N. Y. Saginaw Salt Products Co., Saginaw, Mich.

Calcium Chloride (Anhydrous)

Fales Chemical Co., Inc., Cornwall Landing, N. Y.

Calcium Phosphate

Provident Chemical Wks., St. Louis, Mo.

Calcium Sulphide (Luminous)

Amer. Luminous Products Co., Huntington Park, Calif.

Calcium Stearate

The Synthetic Products Co., Cleveland, Ohio

Camphor

E. J. Barry, New York, N. Y.

Camphor Oil

Magnus, Mabee & Reynard, Inc., New York, N. Y.

Candelilla Wax

Innis, Speiden & Co., Inc., New York, N. Y.

Caramel Color

Alex Fries & Bro., Cincinnati, Ohio

Caraway Oil

Geo. Lueders & Co., New York, N. Y.

Carbolic Oil

Reilly Tar & Chemical Corp., New York, N. Y.

Carbon, Activated

The Jennison-Wright Co., Toledo, Ohio

Carbon Bisulphide

J. T. Baker Chemical Co., Phillipsburg, N. J.

Carbon Black

United Carbon Co., Charleston, W. Va. Binney & Smith, New York, N. Y.

Carbon, Decolorizing

Darco Sales Corp., New York, N. Y.

Carbon Tetrachloride

Niagara Smelting Corp., Niagara Falls, N. Y.

Cardamom Seed

Newmann-Buslee & Wolfe, Inc., Chicago, Ill.

Carnauba Wax

Frank B. Ross Co., Inc., New York, N. Y.

Cannin

The Casein Mfg. Co. of America, Inc., New York, N. Y.

Castile Soap Conti Products Corp., New York, N. Y.

Castor Oil
The Baker Castor Oil Co., New York, N. Y.

Castor Oil, Sulphonated
Jacques Wolf & Co., Passaic, N. J.

Celluloid Corp., New York, N. Y.

Celluloid Scrap Moses Serinsky Co., Indianapolis, Ind.

Cellulose Acetate
Celanese Corp. of America, New York, N. Y.

Cellulose Nitrate
Merrimac Chemical Co., Everett, Mass.

Ceresin Wax Sherwood Petroleum Co., Inc., Brooklyn, N. Y.

Cetyl Alcohol Hummel Chemical Co., Inc., 90 West St., New York, N. Y.

Chalk, Precipitated
Charles B. Chrystal Co., Inc., New York, N. Y.

Charcoal
Chas. L. Read & Co., Inc., New York, N. Y.
Western Charcoal Co., Chicago, Ill.

China Clay
Taintor Trading Co., New York, N. Y.

China Wood Oil
Balfour, Guthrie & Co., Ltd., New York, N. Y.

Chloramine
Abbott Laboratories, No. Chicago, Ill.

Chlorine (Liquid)
Electro Bleaching Gas Co., 9 E. 41st St., New York, N. Y.

Chloroform
The Dow Chemical Co., Midland, Michigan

Chlorophyll
Amer. Chlorophyll, Inc., New York, N. Y.
Pylam Products Co., New York, N. Y.

Cholesterin
Digestive Ferments Co., Detroit, Michigan
Merck & Co., Inc., Rahway, N. J.

Chrome Green Kentucky Color & Chem. Co., Louisville, Ky.

Chrome Yellow Ansbacher-Siegle Corp., Rosebank, N. Y.

Chromic Acid
Mutual Chemical Co. of America, New York, N. Y.

Chromium Oxide
O. Hommel Co., Inc., Pittsburgh, Pa.

Citral Givaudan-Delawanna, Inc., New York, N. Y.

Citric Acid
Chas. Pfizer & Co., Inc., New York, N. Y.

Citronella Oil H. C. Ryland, Inc., New York, N. Y. Clay
Kentucky Clay Mining Co., Mayfield, Ky.
Olive Branch Minerals Co., Cairo, Ill.

Crowley Tar Products Co., New York, N. Y.

Coal Tar Colors
H. Kohnstamm & Co., New York, N. Y.

Cobalt Acetate
Fred L. Brooke Co., Chicago, Ill.

Cobalt Driers
McGean Chemical Co., Cleveland, Ohio

Cobalt Linoleate
The McGean Chemical Co., Cleveland, Ohio

Cocoa Butter
Alpha Lux Co., Inc., New York, N. Y.
Thomas J. Shields Co., New York, N. Y.

Coconut Butter
Procter & Gamble Co., Cincinnati, Ohio

Coconut Oil Franklin Baker Co., Hoboken, N. J.

Coconut Oil Fatty Acid
Aeme Oil Corp., Chicago, Ill.

Cod Liver Oil
H. H. Rosenthal & Co., Inc., New York, N. Y.

Collodion
Charles Cooper & Co., New York, N. Y.

Colors, Dry
Holland Aniline Dye Co., Holland, Mich.

Colors, Oil Soluble Commonwealth Color & Chem. Co., Brooklyn, N. Y.

Copper Carbonate
Chas. Copper & Co., New York, N. Y.
Jungmann & Co., Inc., New York, N. Y.

Copper Cyanide Charles Hardy, Inc., New York, N. Y.

Copper Oxides
The O. Hommel Co., Inc., 209 Fourth Ave., Pittsburgh, Pa.

Copper Sulphate
Barada & Page, Inc., Kansas City, Mo.

Corn Oil
American Maize Products Co., New York, N. Y.

Corn Sugar Staley Sales Corp., Decatur, Ill.

Corn Syrup Clinton Co., Clinton, Ia. Corn Products Refining Co., New York, N. Y.

Cottonseed Oil (Crude)
Battleboro Oil Co., Battleboro, N. C.
Welch, Holme & Clark Co., New York, N. Y.

Coumarin
Maywood Chem. Works, Maywood, N. J.

Coumarone Resin
Barrett Co., New York, N. Y.
Neville Co., Pittsburgh, Pa.

Cream of Tartar

The Harshaw Chemical Co., Cleveland, Ohio

Creosote

Koppers Products Co., Pittsburgh, Pa.

Cresols

Coopers Creek Chem. Co., W. Conshohocken, Pa. Reilly Tar & Chemical Corp., New York, N. Y.

Cresylic Acid

The Barrett Co., New York, N. Y.

Cryolite

Vitro Mfg. Co., Pittsburgh, Pa.

Cyclohexanol

E. I. Du Pont de Nemours Co., Wilmington, Del.

Damar Gum

Geo. H. Lincks, New York, N. Y.

Degras

Amer. Lanolin Corp., Lawrence, Mass.

Derris Extract

Seacoast Laboratories, New York, N. Y.

Derris Root

W. Benkert & Co., Inc., New York, N. Y.

Dextrins

Morningstar, Nicol, Inc., New York, N. Y.

Diastase

Takamine Laboratory, Inc., Clifton, N. J.

Diatomaceous Earth

Dicalite Co., New York, N. Y.

Dibutylphthalate

The Kessler Chemical Corp., New York, N. Y.

Dichlorbenzol

Hooker Electro Chemical Co., New York, N. Y.

Diethyleneglycol

Carbide & Carbon Chemicals Corp., New York, N. Y.

Diethylphthalate

Van Dyk & Co., Inc., Jersey City, N. J.

Digtycol Oleate

Glyco Products Co., Inc., New York, N. Y.

Diglycol Laurate

Glyco Products Co., Inc., New York, N. Y.

Diglycol Stearate

Glyco Products Co., Inc., New York, N. Y.

Dioxan

Carbide & Carbon Chem. Corp., New York, N. Y.

Dipentene

Hercules Powder Co., Wilmington, Del.

Diphenyl

Swann Chemical Co., New York, N. Y.

Drop Black

Wilckes-Martin-Wilckes Co., New York, N. Y.

Dyestuffs

National Aniline & Chemical Co., Inc., New York, N. Y.

Egg, Dried

W. P. Pray, New York, N. Y.

Egg Yolk

Stein, Hall & Co., New York, N. Y.

Ephedrine

Abbott Laboratories, No. Chicago, Ill.

Epsom Salt

General Chemical Co., New York, N. Y.

Essential Oils

Compagnie Duval, New York, N. Y.

Ester Gum

John D. Lewis, Inc., Providence, R. I. Paramet Chemical Corp., Long Island City, N. Y.

Carbide & Carbon Chemicals Corp., New York, N. Y.

Ethyl Acetate

Merrimac Chemical Co., Boston, Mass.

Ethyl Cellulose

Advance Solvents & Chem. Corp., New York, N. Y.

Ethylamine

F. C. Bersworth Labs., Framingham, Mass.

Ethyl Lactate

American Cyanamid & Chemical Corp., New York, N. Y.

Ethylene Diamine

F. C. Bersworth Labs., Framingham, Mass.

Ethylene Dichloride

Dow Chemical Co., Midland, Mich.

Ethyleneglycol

Carbide & Carbon Chemicals Corp., New York, N. Y.

Chas. Fishbeck Co., New York, N. Y.

Feldspar

Consolidated Feldspar Corp., Trenton, N. J.

Fillers

C. K. Williams & Co., Easton, Pa.

Film Scrap

Horn-Jefferys & Co., Burbank, Calif.

Fish Glue

C. B. Hewitt & Bro., New York, N. Y.

Fish Oil

Falk & Co., Pittsburgh, Pa.

Flaxseed

Bisbee Linseed Co., Philadelphia. Pa.

Fluorspar

Hillside Fluor Spar Mines, Chicago, Ill.

Formic Acid Victor Chem. Works, Chicago, Ill.

Formaldehyde

Heyden Chemical Corp., New York, N. Y.

Fuller's Earth

L. A. Salmon & Bro., New York, N. Y. Sinclair Refining Co., Olmstead, Ill.

Fusel Oil

Empire Distilling Corp., New York, N. Y.

Gallic Acid

Eastman Kodak Co., Rochester, N. Y.

Gamboge

Frank B. Ross Co., New York, N. Y.

Gelatin

Atlantic Gelatine Co., Woburn, Mass.

Geraniol

Kay-Fries Chem., Inc., New York, N. Y.

Geranium Lake

Interstate Color Co., Inc., New York, N. Y. R. F. Revson Co., New York, N. Y.

Geranium Oil

Schimmel & Co., New York, N. Y.

Gilsonite

George H. Lincks, New York, N. Y. Utah Gilsonite Co., St. Louis, Mo.

Ginseng

C. H. Lewis & Co., New York, N. Y.

Glandular Products

The Wilson Laboratories, Chicago, Ill.

Glauber Salt

Iowa Soda Products Co., Council Bluffs, Ia.

Clare

Cudahy Packing Co., Chicago, Ill.

Glycerin

Colgate-Palmolive-Peet Co., Chicago, Ill.

Glyceryl Mono Stearate

Glyco Products Co., Inc., New York, N. Y.

Glyceryl Phthalate

Glyco Products Co., Inc., New York, N. Y.

Glyceryl Stearate

Glyco Products Co., Inc., New York, N. Y.

Glycol Oleate

Glyco Products Co., Inc., New York, N. Y.

Glycol Phthalate
Glyco Products

Glyco Products Co., Inc., New York, N. Y.

Glycol Stearate

Glyco Products Co., Inc., New York, N. Y.

Gold Chloride

Mallinckrodt Chemical Works, St. Louis, Mo.

Graphite

Adolphe Hurst & Co., Inc., New York, N. Y. Asbury Graphite Mills, Asbury Park, N. J.

Gum Arabic
T. M. Duche & Sons, New York, N. Y.

Gum Benzoin .

Peek & Velsor, Inc., New York, N. Y.

Gum Copal

George H. Lincks, New York, N. Y.

Gum Damar

Thurston & Braidich, New York, N. Y.

Gum Karaya

Frank-Vliet Co., Inc., New York, N. Y.

Gum, Locust Bean

Innis, Speiden Co., New York, N. Y.

Gum Manila

Stroock & Wittenberg Corp., New York, N. Y.

Gum Tragacanth

E. Meer & Co., Inc., New York, N. Y. J. L. Hopkins & Co., New York, N. Y.

U. S. Phosphoric Prod. Corp., New York, N. Y.

Hemlock Bark

Tanners Supply Co., Grand Rapids, Mich.

Henna Leaves

S. B. Penick & Co., New York, N. Y.

Herbs

John Clarke & Co., New York, N. Y.

Hexamethylenetetramine

Heyden Chemical Corp., New York, N. Y.

Hydrochloric Acid

General Chemical Co., New York, N. Y.

Hydrogen Peroxide

The Warner Chemical Co., New York, N. Y.

Hydroquinone

Eastman Kodak Co., Rochester, N. Y.

Ichthyol

Merck & Co., Rahway, N. J.

Indigo

L. E. Ransom Co., New York, N. Y.

Indium

Belmont Smelting & Refining Works, Brooklyn, N. Y.

Invert Sugar

Nulomoline Co., New York, N. Y.

Iodine

New York Quinine & Chemical Wks., Inc., Brooklyn, N. Y.

Iridium

Baker & Co., Inc., Newark, N. J.

Irish Moss

S. B. Penick & Co., New York, N. Y.

Iron Ammonium Citrate

Schuykill Chem. Co., Philadelphia, Pa.

Iron Chloride

Chicago Copper & Chem. Co., Blue Island, Ill.

Iron Oxide

Binney & Smith Co., New York, N. Y.

Isopropyl Acetate

A. K. Hamilton, New York, N. Y.

Isopropyl Alcohol

Carbide & Carbon Chemicals Corp., New York, N. Y.

Insect Wax, Chinese

Frank B. Ross Co., Inc., New York, N. Y.

Ivory Black

Binney & Smith Co., New York, N. Y.

Japan Wax

Smith & Nichols, Inc., New York, N. Y.

Kerosene

Colonial Beacon Oil Co., Everett, Mass.

Kerosene, Deodorized

Sherwood Petroleum Co., Brooklyn, N. Y.

Laboratory Equipment

Central Scientific Co., Chicago, Ill. Chemical Publ. Co. of N. Y., Inc., New York, N. Y. Chicago Apparatus Co., Chicago, I Eimer & Amend, New York, N. Y.

Experimenter's Supply Co., New York, N. Y.

Fisher Scientific Co., Pittsburgh, Pa.

N. J. Laboratory Supply Co., Newark, N. J. Scientific Glass Apparatus Co., Bloomfield, N. J.

Maas & Waldstein, Newark, N. J.

Lactic Acid

Apex Chemical Co., Inc., New York, N. Y.

Lamp Black

Binney & Smith Co., New York, N. Y. L. Martin Co., New York, N. Y.

Lanolin

American Lanolin Corp., Lawrence, Mass.

Merck & Co., Inc., Rahway, N. J. Pfaltz & Bauer, New York, N. Y.

Lard Oil

Enterprise Animal Oil Co., Philadelphia, Pa.

Lauryl Alcohol and Sulphonate

E. I. Du Pont de Nemours & Co., Wilmington, Del.

Lavender Oil

Van Ameringen-Haebler, Inc., New York, N. Y.

Lead Acetate

National Lead Co., New York, N. Y.

Lead Arsenate

Barada & Page, Inc., Kansas City, Mo. General Chemical Co., New York, N. Y.

Lead and Its Oxides

The Eagle-Picher Sales Co, Cincinnati, Ohio

Lecithin

American Lecithin Corp., New York, N. Y.

Lemon Juice, Concentrated

Mutual Citrus Products Co., Anaheim, Calif.

Lemon Oil

D. W. Hutchinson & Co., Inc., New York, N. Y.

MacAndrews & Forbes Co., New York, N. Y.

Lime

J. E. Baker Co., York, Pa. Chazy Marble Lime Co., Inc., Chazy, N. Y.

Limestone

F. E. Schundler & Co., Joliet, Ill.

Linoleic Acid

Glyco Products Co., Inc., New York, N. Y.

Linseed Oil

Bisbee Linseed Co., Philadelphia, Pa.

Litharge

The Eagle-Picher Lead Co., Cincinnati, Ohio

Lithopone

Krebs Pigment & Color Corp., Newark, N. J. Marshall Dill Co., San Francisco, Calif.

Locust Bean Powder

T. M. Duche & Sons, New York, N. Y.

Logwood Extract

American Dyewood Co., New York, N. Y.

Lycopodium

McKesson & Robbins, Inc., New York, N. Y.

Magnesia

Philip Carey Co., Lockland, O.

Magnesite

General Magnesite & Magnesia Co., Philadelphia, Pa.

Magnesium Carbonate

Merck & Co., Inc., Rahway, N. J.

Magnesium Chloride

Wishnick-Tumpeer, Inc., New York, N. Y.

Magnesium Powder

Belmont Smelting & Refining Wks., Inc., Brooklyn, N. Y.

Maleic Acid

Nat'l Aniline & Chem. Wks., New York, N. Y.

Manganese

Ajax Metal Co., Philadelphia, Pa.

Marble Dust

Hammil & Gillespie, Inc., New York, N. Y.

Manganese Dioxide

B. F. Drakenfeld & Co., Inc., New York, N. Y.

Menhaden Oil

Robert Badcock & Co., New York, N. Y.

Menthol

Chas. L. Huisking & Co., Inc., New York, N. Y.

Mercury

Chas. L. Huisking & Co., Inc., New York, N. Y.

George Uhe Co., New York, N. Y.

Methanol

Wm. S. Gray & Co., New York, N. Y.

Methyl Acetate

Carbide & Carbon Chem. Corp., New York, N. Y.

Methyl Acetone

Delta Chem. & Iron Co., Wells, Mich.

Methyl Anthranilate

Florasynth Laboratories, New York, N. Y.

Methyl p-Hydroxybenzoate

Heyden Chemical Corp., New York, N. Y.

Methyl Salicylate

Dow Chemical Co., Midland, Michigan

Mica

Southern Mica Co., Franklin, N. C.

Milk Sugar

Mallinckrodt Chemical Wks., St. Louis, Mo.

Mineral Rubber

Barber Asphalt Co., Philadelphia, Pa.

Mineral Spirits

Amer. Mineral Spirit Co., New York, N. Y.

Montan Wax

Strahl & Pitsch, New York, N. Y.

Naphtha

Deep Rock Oil Corp., Chicago, Ill.

Naphthalene

The Barrett Co., New York, N. Y.

Naphthenic Acid

Glyco Products Co., Inc., New York, N. Y.

National Oil Products Co., Harrison, N. J.

Nickel Chloride

Chas. Cooper & Co., New York, N. Y.

Nickel Sulphate

The Harshaw Chemical Co., Cleveland, O.

Nicotine

Tobacco By-Products & Chemical Corp., Louisville, Ky.

Nicotine Sulphate

Lattimer-Goodwin Chemical Co., Grand Junction, Colo.

Nitre Cake

Trojan Powder Co., Allentown, Pa.

Nitric Acid

Monsanto Chemical Co., St. Louis, Mo.

Nitrobenzol

Calco Chem. Co., Bound Brook, N. J.

Nitrocellulose

E. I. Du Pont de Nemours & Co., Inc., Parlin, N. J.

Smith Chemical & Color Co., Brooklyn, N. Y.

Oil, Citronella

D. W. Hutchinson & Co., Inc., New York, N. Y.

Oil, Mineral

Standard Oil Co. of California, San Francisco, Calif.

Oil, Olive

Leghorn Trading Co., Inc., New York, N. Y.

Oiticica Oil

L. N. Jackson & Co., New York, N. Y.

Century Stearic Acid Wks., New York, N. Y.

Oleoresins

Seeley & Co., New York, N. Y.

Olive Oil, Sulphonated

Royce Chem. Co., Carlton Hill, N. J.

Orange Oil

Dodge & Olcott Co., New York, N. Y.

Ortho Dichlorbenzene

Hooker Electrochemical Co., New York, N. Y.

Oxalic Acid

Mutual Chemical Co. of America, New York, N. Y.

Oxgall

Wilson Labs., Chicago, Ill.

Oxyger

Cheney Chemical Co., Cleveland, O.

Oxyguinoline Sulphate

Benzol Products Co., Newark, N. J.

Ozokerite Wax

Strohmeyer & Arpe Co., New York, N. Y.

Palm Kernel Oil

Franklin Baker Co., Hoboken, N. J.

Palm Oil

Wishnick-Tumpeer, Inc., New York, N. Y.

Paraffin Oils

S. Schwabacher & Co., Inc., New York, N. Y.

Paraffin Wax

Oil States Petroleum Co., New York, N. Y.

Paraldehyde

Heyden Chem. Corp., New York, N. Y.

Para Aminophenol

Verona Chem Co., Newark, N. J.

Para-Phenylenediamine

Amido Products Co., New York, N. Y.

Paris White

Southwark Mfg. Co., Camden, N. J.

Peanut Oil

Elbert & Co., New York, N. Y.

Pearl Essence

Mearl Corp., New York, N. Y.

Pectin.

Calif. Fruit Growers' Exchange, Ontario, Calif.

Peppermint Oil

Magnus, Mabee & Reynard, Inc., New York, N. Y. The Sparhawk Co., Sparkhill, N. Y.

Perilla Oil

S. L. Jones & Co., San Francisco, Calif.

Petrolatum

Pennsylvania Refining Co., Butler, Pa.

Petroleum Jelly

L. Sonneborn Sons, Inc., New York, N. Y.

Petroleum Spirits

Sun Oil Co., Philadelphia, Pa.

Phenol

American-British Chemical Supplies, Inc., New York, N. Y.

Phenol-Formaldehyde Resins

Durite Plastics, Philadelphia, Pa.

Phosphoric Acid

Victor Chemical Works, Chicago, Ill.

Phosphorus

International Selling Corp., New York, N. Y.

Phthalic Anhydride

Monsanto Chem. Co., St. Louis, Mo.

Pine Oil

General Naval Stores Co., Inc., New York, N. Y.

Pine Tar

Southern Pine Chem. Co., Jacksonville, Fla.

Pitch

Robert Rauh, Inc., Newark, N. J.

Plaster of Paris

Whittaker, Clark & Daniels, Inc., New York, N. Y.

Potash, Caustic

Niagara Alkali Co., New York, N. Y.

Potassium Carbonate

Joseph Turner & Co., New York, N. Y.

Potassium Chlorate

Joseph Turner & Co., New York, N. Y.

Potassium Hydroxide

Merck & Co., Inc., Rahway, N. J.

 ${\it Potassium~Iodide}$

New York Quinine & Chemical Wks., Inc., Brooklyn, N. Y.

Potassium Oleate

Glyco Products Co., Inc., New York, N. Y. Carl F. Miller & Co., Seattle, Washington

Potassium Permanganate

Carus Chemical Co., Inc., La Salle, Ill.

Potassium Silicate

Philadelphia Quartz Co., Philadelphia, Pa.

Prussian Blue Kentucky Color & Chem. Co., Louisville, Ky.

Charles B. Crystal Co., New York, N. Y.

Psyllium Seeds

Laxseed Co., New York, N. Y.

Pyrethrum Extract

McLaughlin, Gormley, King & Co., Minneapolis, Minn.

Pyrethrum

S. B. Penick & Co., New York, N. Y.

Pyrogallic Acid

Zinsser & Co., Inc., Hastings-on-Hudson, N. Y.

Pyroxylin Solutions

Egyptian Lacquer, Kearney, N. J.

Quince Seed

J. L. Hopkins & Co., New York, N. Y.

Quinine Bisulphate

R. W. Greef & Co., Inc., New York, N. Y.

Rapeseed Oil

Balfour, Guthrie & Co., Ltd., New York, N. Y.

Red Oil

Century Stearic Acid Candle Wks., New York, N. Y.

Resins, Synthetic

Beck, Koller & Co., Inc., Detroit, Mich. Marshall Dill, San Francisco, Calif.

Resorcin

Penn. Coal Products Co., Petrolia, Pa.

Rhodium

Baker & Co., Inc., Newark, N. J.

Rochelle Salts

Chas. Pfizer & Co., Inc., New York, N. Y.

Rose Water

Geo. Lueders & Co., New York, N. Y.

Rosin

General Naval Stores Co., Inc., New York, N. Y.

Rosin Oil

National Rosin Oil & Size Co., New York, N. Y.

Rotenone

Thorocide, Inc., St. Louis, Mo.

Rubber

Earle Bros., New York, N. Y.

Rubber Latex

Littlejohn & Co., Inc., New York, N. Y.

Saccharine

Heyden Chemical Corp., New York, N. Y.

Salicylic Acid

The Dow Chemical Co., Midland, Mich.

Sal Soda

Church & Dwight Co., Inc., New York, N. Y.

Salt

Morton Salt Co., Chicago, Ill.

Salt Cake

Amer. Cyanamid & Chem. Corp., New York, N. Y.

Saltpetre

Croton Chem. Corp., Brooklyn, N. Y.

San on in

Experimenters Supply Co., New York, N. Y. Jungmann & Co., New York, N. Y.

Selenium

Amer. Metal Co., New York, N. Y.

Shellac

Wm. Zinsser & Co., New York, N. Y.

Shellac Wax

Adolphe Hurst & Co., New York, N. Y.

Siennas

Fezandie & Sperrie, Inc., New York, N. Y.

Silica

Barnsdall Tripoli Corp., Seneca, Mo.

Silne

Handy & Harman, New York, N. Y.

пациу

Silver Cyanide Chas. Cooper & Co., New York, N. Y.

Silver Nitrate

Eastman Kodak Co., Rochester, N. Y.

Soda Ash

Diamond Alkali Co., Pittsburgh, Pa.

Soda, Caustic

Mathieson Alkali Works, Inc., New York, N. Y.

Soda, Sal

Consolidated Chem. Sales Corp., Newark, N. J.

Sodium Aluminate

National Aluminate Corp., Chicago, Ill.

Sodium Arsenite

Harrison Mfg. Co., Rahway, N. J.

Sodium Benzoate

Hooker Electrochemical Co., New York, N. Y.

Sodium Bicarbonate

Church & Dwight Co., Inc., New York, N. Y.

Sodium Bichromate

Prior Chem. Corp., New York, N. Y.

Sodium Bisulphite

The Grasselli Chemical Co., Cleveland, Ohio

Sodium Carbonate

Solvay Sales Corporation, New York, N. Y.

Sodium Choleate

Difco Laboratories, Inc., Detroit, Mich.

Sodium Cyanide

E. I. Du Pont de Nemours & Co., Inc., Wilmington, Del.

Sodium Fluoride

American Cyanamid & Chemical Corp., New York, N. Y.

Sodium Hydrosulphite

Royce Chemical Co., Carlton Hill, N. J.

Sodium Hydroxide

Merck & Co., Inc., Rahway, M. J.

Sodium Hypochlorite

Delta Chemical Mfg. Co., Baltimore, Md. Mathieson Alkali Wks., Inc., New York, N. Y.

Sodium Hypochlorite Liquid Riverside Chemical Co., No. Tonawanda, N. Y.

Sodium Hyposulphite

The Grasselli Chemical Co., Cleveland, Ohio

Sodium Metaphosphate

Buromin Co., Pittsburgh, Pa.

Sodium Metasilicate

Philadelphia Quartz Co., Philadelphia, Pa.

Sodium Nitrate

Battelle & Renwick, New York, N. Y.

Sodium Nitrite

Solvay Sales Corp., New York, N. Y.

Sodium Perborate

E. I. Du Pont de Nemours & Co., Inc., Wilmington, Del.

Sodium Phosphate

Swann Chemical Co., New York, N. Y.

Sodium Resinate

Paper Makers Chem. Corp., Wilmington, Del.

Sodium Silicate

Mechling Bros. Chemical Co., Camden, N. J. Philadelphia Quartz Co., Philadelphia, Pa. Standard Silicate Co., Pittsburgh, Pa.

Sodium Silico Fluoride
The Grasselli Co., Cleveland, Ohio

Sodium Sulphate
General Chem. Co., New York, N. Y.

Sodium Stannate
Harshaw Chem. Co., Cleveland, Ohio

Sodium Sulphite
Mechling Bros. Chemical Co., Camden, N. J.

Sodium Tungstate
J. T. Baker Chem. Co., Phillipsburg, N. J.

Solvent Naphtha Barrett Co., New York, N. Y.

Sorbitol
Atlas Powder Co., Wilmington, Del.

Soybean Oil Spencer Kellogg & Sons Sales Corp., Buffalo, N. Y. Arthur C. Trask Co., Chicago, Ill.

Sperm Oil Cook Swan Co., Inc., New York, N. Y.

Spermaceti Strahl & Pitsch, New York, N. Y.

Squill
S. B. Penick & Co., New York, N. Y.

Starch
Starch Products Co., New York, N. Y.

Stearic Acid Candle Wks., New York, N. Y.

Stearin M. Werk Co., Cincinnati, Ohio

Stearine Pitch
A. Gross & Co., New York, N. Y.

Strontium Nitrate
Grasselli Chem. Co., Cleveland, Ohio

Strychnine Chas. Pfizer & Co., New York, N. Y.

Sulphonated Castor Oil Burkard-Schier Chem. Co., Chattanooga, Tenn.

Sulphonated Olive Oil Jacques Wolf & Co., Passaic, N. J.

Sulphur Stauffer Chemical Co. of Texas, Freeport, Tex.

Sulphur Dioxide
Virginia Smelting Co., Boston, Mass.

Sulphuric Acid
Merrimac Chemical Co., Everett Sta., Boston, Mass.

Talc
Charles B. Crystal Co., Inc., New York, N. Y.

Tallow

Welch, Holme & Clark Co., Inc., New York, N. Y.

Tartaric Acid

R. W. Greeff & Co., Inc., New York, N. Y.

Tar Acid Oil

Barrett Co., New York, N. Y.

Tartar Emetic

Apex Chem. Co., New York, N. Y.

Tea Seed Oil

Lundt & Co., New York, N. Y.

D. W. Hutchinson & Co., New York, N. Y.

Tetrachlorethane

Dow Chemical Co., Midland, Mich.

Tetrachlorethylene

E. I. Du Pont de Nemours & Co., Wilmington, Del.

Thallium Sulphate

Jungmann & Co., Inc., New York, N. Y.

Thio carbamilid

Monsanto Chemical Co., St. Louis, Mo.

Jungmann & Co., New York, N. Y.

Thymol

Sherka Chemical Co., Inc., Bloomfield, N. J.

Union Smelting & Refining Co., Inc., Newark, N. J.

Tin Chloride

Seldner & Enequist, Inc., Brooklyn, N. Y.

Tin Oxide

McGean Chemical Co., Cleveland, Ohio

Parke, Davis & Co., Detroit, Mich.

Titanium Dioxide

Marshall Dill, San Francisco, Calif. R. T. Vanderbilt Co., New York, N. Y.

Jones & Laughlin Steel Corp., Pittsburgh, Pa.

Niacet Chemicals Corp., Niagara Falls, N. Y.

Tricresyl Phosphate

R. W. Greeff & Co., Inc., New York, N. Y.

Triethanolamine

Experimenter's Supply Co. (small lots), New York, N. Y. Carbide & Carbon Chem. Co. (large lots), New York, N. Y.

Triethanolamine Oleate

Glyco Products Co., Inc., New York, N. Y.

Marshall Dill Co., San Francisco, Calif.

Triethanolamine Stearate

Glyco Products Co., Inc., New York, N. Y. Carl F. Miller & Co., Seattle, Washington

Triphenylquanadine

E. I. Du Pont de Nemours & Co., Wilmington, Del.

Triphenylphosphate
Monsanto Chemical Co., St. Louis, Mo.

Tamms Silica Co., Chicago, Ill.

Tungsten
Fansteel Products Co., No. Chicago, Ill.

Turkey Red Oil
National Oil Products Co., Inc., Harrison, N. J.

Antwerp Naval Stores Co., Inc., Boston, Mass. General Naval Stores Co., New York, N. Y.

Turpentine Substitute
Anderson-Prichard Oil Corp., Oklahoma City, Okla.

Turpentine (Venice) National Rosin Oil & Size Co., New York, N. Y.

Turtle Oil Edwin Seebach Co., New York, N. Y.

Ultramarine Blue Standard Ultramarine Co., Huntington, W. Va.

Fezandie & Sperrle, Inc., New York, N. Y.

Uranium Nitrate
Harshaw Chemical Co., Cleveland, Ohio

Sherka Chemical Co., Inc., Bloomfield, N. J.

Vanilla Beans
Thurston & Braidich, New York, N. Y.

Vanillin
Seeley & Co., Inc., New York, N. Y.

Van Ameringen-Haebler, Inc., New York, N. Y. Varnish Gums and Resins

Amer. Cyanamid & Chem. Corp., New York, N. Y.

Vat Colors
Amer. Aniline Products, Inc., New York, N. Y.

Vegetable Colors
L. E. Ransom Co., New York, N. Y.

Vermiculite
Hill Bros. Chem. Co., Los Angeles, Calif.

Vermilion Fezandié & Sperrlé, Inc., New York, N. Y.

Vinyl Acetate
Niagara Chemicals Corp., Niagara Falls, N. Y.

Vinyl Chloride Carbide & Carbons Chem. Corp., New York, N. Y.

Wax, Synthetic Glyco Products Co., Inc., New York, N. Y.

Wetting Out Agents
Glyco Products Co., Inc., New York, N. Y.

Whiting
Columbia Alkali Corp., New York, N. Y.
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Witch Hazel Extract E. E. Dickinson Co., Essex, Conn. White Arsenic

H. H. Rosenthal Co., New York, N. Y.

White Lead

National Lead Co., New York, N. Y.

Wood Flour

D. H. Litter Co., New York, N. Y. Wood Flour, Inc., Manchester, N. H.

Xylol

The Barrett Co., New York, N. Y.

Yeast

Standard Brands, Inc., New York, N. Y.

Zinc

Hegeler Zinc Co., Danville, Ill.

Zinc Carbonate

Wishnick-Tumpeer, Inc., New York, N. Y.

Zinc Chloride

Wishnick-Tumpeer, Inc., New York, N. Y.

Zinc Chromate

E. M. & F. Waldo, Inc., Muirkirk, Md.

Zinc Oxide

Merck & Co., Inc., Rahway, N. J. N. J. Zinc Co., New York, N. Y.

Zinc Stearate

Merck & Co., Inc., Rahway, N. J. Wishnick-Tumpeer, Inc., New York, N. Y.

Zinc Sulphate

W. R. Russell & Co., New York, N. Y. Virginia Smelting Co., West Norfolk, Va.

Zirconium Oxide

Foote Mineral Co., Philadelphia, Pa.

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Canadian Industries, Ltd., Toronto
Chas. Tennant & Co. (Canada), Ltd., 372 Bay St., Toronto
Merck & Co., Ltd., Montreal and Toronto
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British Drug Houses (Canada), Ltd., Terminal Warehouses, Toronto
Shanahan Chemicals, Ltd., Ft. of Campbell Ave., Vancouver, B. C.
Chemicals, Ltd., 384 St. Paul's St. W., Montreal

India

Purshotamdass Popatlal & Co., 37 Hamam St., Fort, Bombay Imperial Chemical Industries (India), Ltd., Imperial Chemical House, Ballard Estate, Bombay Ciba (India), Ltd., Post Box 479, Bombay

Holland.

N. V. Chemische Fabriek Servo, Delden (Twente), Holland W. A. Scholten's Chemical Works, Ltd., Groningen, Holland

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27 Chesser St., Adelaide
414 Kent Ave., Sidney

19 Lower Tory St., Wellington, New Zealand

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W. H. Goetz, Calle Sarandi 315, Buenos Aires

J. M. Sierra, Aquiar 73 Dpt. 710, Apartado 363. Havana

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